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<p>(21) International Application Number: PCT/US97/04460 (22) International Filing Date: 26 March 1997 (26.03.97) (30) Priority Data: 60/014,288 29 March 1996 (29.03.96) US (60) Parent Application or Grant (63) Related by Continuation US 60/014,288 (CON) Filed on 29 March 1996 (29.03.96) (71) Applicant (for all designated States except US): G.D. SEARLE & CO. [US/US]; Corporate Patent Dept., P.O. Box 5110, Chicago, IL 60680-5110 (US). (72) Inventors; and (75) Inventors/Applicants (for US only): CHEN, Barbara, B. [US/US]; 1921 Robincrest Lane, Glenview, IL 60025 (US). CHEN, Helen, Y. [US/US]; 7 Baldwin Terrace, Livingston, NJ 07039 (US). GESICKI, Glen, J. [US/US]; 6837 Davis Street, Morton Grove, IL 60053 (US). HAACK,</p>	<p>Richard, A. [US/US]; 5356 North Luna, Chicago, IL 60030 (US). MALECHA, James, W. [US/US]; 1121 Tracy Lane, Libertyville, IL 60048 (US). PENNING, Thomas, D. [US/US]; 374 Larch, Elmhurst, IL 60136 (US). RICO, Joseph, G. [US/US]; 524 Weatherby Terrace Drive, Ballwin, MO 63021 (US). ROGERS, Thomas, E. [US/US]; 755 Trago Creek Drive, Ballwin, MO 63021 (US). RUMINSKI, Peter, G. [US/US]; 391 Crestbury, Ballwin, MO 63011 (US). RUSSELL, Mark, A. [GB/US]; 475 Cross Road, Gurnee, IL 60031 (US). YU, Stella, S. [US/US]; 7801 Maple Street, Morton Grove, IL 60053 (US). (74) Agents: KOVACEVIC, Cynthia, S. et al.; G.D. Searle & Co., Corporate Patent Dept., P.O. Box 5110, Chicago, IL 60680-5110 (US). (81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ARIPO patent (GH, KE, LS, MW, SD, SZ, UG), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). Published With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</p>	
<p>(54) Title: PARA-SUBSTITUTED PHENYLPROPANOIC ACID DERIVATIVES AS INTEGRIN ANTAGONISTS</p> <div style="text-align: center;"> <p>(I)</p> </div> <p>(57) Abstract</p> <p>The present invention relates to a class of compounds represented by Formula (I) or a pharmaceutically acceptable salt thereof, pharmaceutical composition comprising compounds of Formula (I), and methods of selectively inhibiting or antagonizing the $\alpha_v\beta_3$ integrin.</p>		

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PARA-SUBSTITUTED PHENYLPROPANOIC ACID DERIVATIVES AS INTEGRIN ANTAGONISTS

Field of the Invention

5 The present invention relates to pharmaceutical agents (compounds) which are useful as $\alpha_v\beta_3$ integrin antagonists and as such are useful in pharmaceutical compositions and in methods for treating conditions mediated by $\alpha_v\beta_3$ by inhibiting or antagonizing $\alpha_v\beta_3$ integrins.

10

Background of the Invention

Integrins are a group of cell surface glycoproteins which mediate cell adhesion and therefore are useful mediators of cell adhesion interactions
15 which occur during various biological processes. Integrins are heterodimers composed of noncovalently linked α and β polypeptide subunits. Currently eleven different α subunits have been identified and six different β subunits have been identified. The various
20 α subunits can combine with various β subunits to form distinct integrins.

The integrin identified as $\alpha_v\beta_3$ (also known as the vitronectin receptor) has been identified as an integrin which plays a role in various conditions or
25 disease states including tumor metastasis, solid tumor growth (neoplasia), osteoporosis, Paget's disease, humoral hypercalcemia of malignancy, angiogenesis, including tumor angiogenesis, retinopathy, arthritis, including rheumatoid arthritis, periodontal disease, psoriasis and smooth muscle cell migration (e.g.
30 restenosis). Additionally, it has been found that such agents would be useful as antivirals, antifungals and antimicrobials. Thus, compounds which selectively inhibit or antagonize $\alpha_v\beta_3$ would be beneficial for
35 treating such conditions.

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It has been shown that the $\alpha_v\beta_3$ integrin and other α_v containing integrins bind to a number of Arg-Gly-Asp (RGD) containing matrix macromolecules. Compounds containing the RGD sequence mimic extracellular matrix ligands so as to bind to cell surface receptors. However, it is also known that RGD peptides in general are non-selective for RGD dependent integrins. For example, most RGD peptides which bind to $\alpha_v\beta_3$ also bind to $\alpha_v\beta_5$, $\alpha_v\beta_1$ and $\alpha_{IIb}\beta_3$. Antagonism of platelet $\alpha_{IIb}\beta_3$ (also known as the fibrinogen receptor) is known to block platelet aggregation in humans. In order to avoid bleeding side-effects when treating the conditions or disease states associated with the integrin $\alpha_v\beta_3$, it would be beneficial to develop compounds which are selective antagonists of $\alpha_v\beta_3$ as opposed to $\alpha_{IIb}\beta_3$.

Tumor cell invasion occurs by a three step process: 1) tumor cell attachment to extracellular matrix; 2) proteolytic dissolution of the matrix; and 3) movement of the cells through the dissolved barrier. This process can occur repeatedly and can result in metastases at sites distant from the original tumor.

Seftor et al. (Proc. Natl. Acad. Sci. USA, Vol. 89 (1992) 1557-1561) have shown that the $\alpha_v\beta_3$ integrin has a biological function in melanoma cell invasion. Montgomery et al., (Proc. Natl. Acad. Sci. USA, Vol. 91 (1994) 8856-60) have demonstrated that the integrin $\alpha_v\beta_3$ expressed on human melanoma cells promotes a survival signal, protecting the cells from apoptosis. Mediation of the tumor cell metastatic pathway by interference with the $\alpha_v\beta_3$ integrin cell adhesion receptor to impede tumor metastasis would be beneficial.

Brooks et al. (Cell, Vol. 79 (1994) 1157-1164) have demonstrated that antagonists of $\alpha_v\beta_3$ provide a therapeutic approach for the treatment of neoplasia (inhibition of solid tumor growth) since systemic administration of $\alpha_v\beta_3$ antagonists causes dramatic

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regression of various histologically distinct human tumors.

5 The adhesion receptor integrin $\alpha_v\beta_3$ was identified as a marker of angiogenic blood vessels in chick and man and therefore such receptor plays a critical role in angiogenesis or neovascularization. Angiogenesis is characterized by the invasion, migration and proliferation of smooth muscle and endothelial cells. Antagonists of $\alpha_v\beta_3$ inhibit this process by selectively
10 promoting apoptosis of cells in neovasculature. The growth of new blood vessels, or angiogenesis, also contributes to pathological conditions such as diabetic retinopathy (Adonis et al., Amer. J. Ophthalm., Vol. 118, (1994) 445-450) and rheumatoid arthritis (Peacock et al., J. Exp. Med., Vol. 175, (1992), 1135-1138).
15 Therefore, $\alpha_v\beta_3$ antagonists would be useful therapeutic targets for treating such conditions associated with neovascularization (Brooks et al., Science, Vol. 264, (1994), 569-571).

20 It has been reported that the cell surface receptor $\alpha_v\beta_3$ is the major integrin on osteoclasts responsible for attachment to bone. Osteoclasts cause bone resorption and when such bone resorbing activity exceeds bone forming activity it results in
25 osteoporosis (a loss of bone), which leads to an increased number of bone fractures, incapacitation and increased mortality. Antagonists of $\alpha_v\beta_3$ have been shown to be potent inhibitors of osteoclastic activity both *in vitro* [Sato et al., J. Cell. Biol., Vol. 111
30 (1990) 1713-1723] and *in vivo* [Fisher et al., Endocrinology, Vol. 132 (1993) 1411-1413]. Antagonism of $\alpha_v\beta_3$ leads to decreased bone resorption and therefore restores a normal balance of bone forming and resorbing activity. Thus it would be beneficial to provide
35 antagonists of osteoclast $\alpha_v\beta_3$ which are effective inhibitors of bone resorption and therefore are useful in the treatment or prevention of osteoporosis.

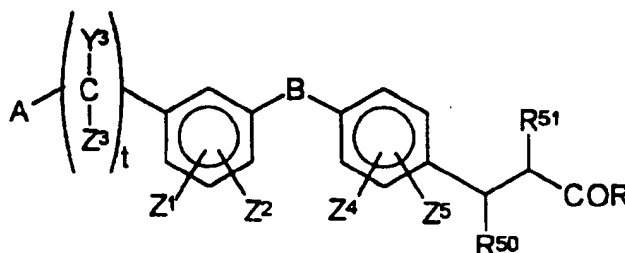
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The role of the $\alpha_v\beta_3$ integrin in smooth muscle cell migration also makes it a therapeutic target for prevention or inhibition of neointimal hyperplasia which is a leading cause of restenosis after vascular procedures (Choi et al., J. Vasc. Surg. Vol. 19(1) (1994) 125-34). Prevention or inhibition of neointimal hyperplasia by pharmaceutical agents to prevent or inhibit restenosis would be beneficial.

White (Current Biology, Vol. 3(9)(1993) 596-599) has reported that adenovirus uses $\alpha_v\beta_3$ for entering host cells. The integrin appears to be required for endocytosis of the virus particle and may be required for penetration of the viral genome into the host cell cytoplasm. Thus compounds which inhibit $\alpha_v\beta_3$ would find usefulness as antiviral agents.

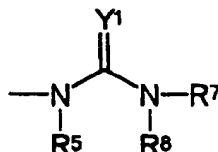
Summary of the Invention

The present invention relates to a class of compounds represented by the Formula I



or a pharmaceutically acceptable salt thereof, wherein

A is



- 5 -

wherein Y^1 is selected from the group consisting of N- R^2 , O, and S;

- 5 R^2 is selected from the group consisting of H; alkyl; aryl; hydroxy; alkoxy; cyano; nitro; amino; aminocarbonyl; alkenyl; alkynyl; alkyl optionally substituted with one or more substituent selected from lower alkyl, halogen, hydroxyl, haloalkyl, cyano, nitro, carboxyl, amino, alkoxy, aryl or
- 10 aryl optionally substituted with one or more halogen, haloalkyl, lower alkyl, alkoxy, cyano, alkylsulfonyl, alkylthio, nitro, carboxyl, amino, hydroxyl, sulfonic acid, sulfonamide, aryl, fused aryl, monocyclic heterocycles, or fused monocyclic
- 15 heterocycles; aryl optionally substituted with one or more substituent selected from halogen, haloalkyl, hydroxy, lower alkyl, alkoxy, methylenedioxy, ethylenedioxy, cyano, nitro, alkylthio, alkylsulfonyl, sulfonic acid,
- 20 sulfonamide, carboxyl derivatives, amino, aryl, fused aryl, monocyclic heterocycles and fused monocyclic heterocycle; monocyclic heterocycles; and monocyclic heterocycles optionally substituted with one or more substituent selected from
- 25 halogen, haloalkyl, lower alkyl, alkoxy, amino, nitro, hydroxy, carboxyl derivatives, cyano, alkylthio, alkylsulfonyl, sulfonic acid, sulfonamide, aryl or fused aryl; or
- 30 R^2 taken together with R^7 forms a 4-12 membered dinitrogen containing heterocycle optionally substituted with one or more substituent selected from the group consisting of lower alkyl, hydroxy and phenyl and wherein said ring optionally
- 35 contains a heteroatom selected from the group consisting of O and S;

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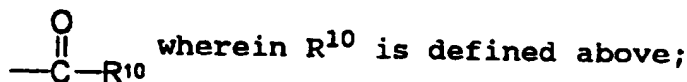
or R^2 taken together with R^7 forms a 5 membered heteroaromatic ring optionally substituted with amino;

5 or R^2 taken together with R^7 forms a 5 membered heteroaromatic ring fused with a phenyl group;

R^7 (when not taken together with R^2) and R^8 are independently selected from the group consisting of H; alkyl; alkenyl; alkynyl; aralkyl; cycloalkyl; bicycloalkyl; aryl; acyl; benzoyl; alkyl optionally substituted with one or more substituent selected from lower alkyl, halogen, hydroxy, haloalkyl, cyano, nitro, carboxyl derivatives, amino, alkoxy, thio, alkylthio, sulfonyl, aryl, aralkyl, aryl optionally substituted with one or more substituent selected from halogen, haloalkyl, lower alkyl, alkoxy, methylenedioxy, ethylenedioxy, alkylthio, haloalkylthio, thio, hydroxy, cyano, nitro, carboxyl derivatives, aryloxy, amido, acylamino, amino, alkylamino, dialkylamino, trifluoroalkoxy, trifluoromethyl, sulfonyl, alkylsulfonyl, haloalkylsulfonyl, sulfonic acid, sulfonamide, aryl, fused aryl, monocyclic heterocycles, fused monocyclic heterocycles; aryl optionally substituted with one or more substituent selected from halogen, haloalkyl, lower alkyl, alkoxy, methylenedioxy, ethylenedioxy, alkylthio, haloalkylthio, thio, hydroxy, cyano, nitro, carboxyl derivatives, aryloxy, amido, acylamino, amino, alkylamino, dialkylamino, trifluoroalkoxy, trifluoromethylsulfonyl, alkylsulfonyl, sulfonic acid, sulfonamide, aryl, fused aryl, monocyclic heterocycles, or fused monocyclic heterocycles; monocyclic heterocycles; monocyclic heterocycles optionally substituted with one or more

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substituent selected from halogen, haloalkyl, lower alkyl, alkoxy, aryloxy, amino, nitro, hydroxy, carboxyl derivatives, cyano, alkylthio, alkylsulfonyl, aryl, fused aryl; monocyclic and bicyclic heterocyclicalkyls; $-\text{SO}_2\text{R}^{10}$ wherein R^{10} is selected from the group consisting of alkyl, aryl and monocyclic heterocycles, all optionally substituted with one or more substituent selected from the group consisting of halogen, haloalkyl, alkyl, alkoxy, cyano, nitro, amino, acylamino, trifluoroalkyl, amido, alkylaminosulfonyl, alkylsulfonyl, alkylsulfonylamino, alkylamino, dialkylamino, trifluoromethylthio, trifluoroalkoxy, trifluoromethylsulfonyl, aryl, aryloxy, thio, alkylthio, and monocyclic heterocycles; and

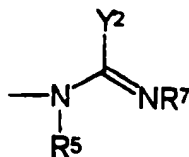


or NR^7 and R^8 taken together form a 4-12 membered mononitrogen containing monocyclic or bicyclic ring optionally substituted with one or more substituent selected from lower alkyl, carboxyl derivatives, aryl or hydroxy and wherein said ring optionally contains a heteroatom selected from the group consisting of O, N and S;

R^5 is selected from the group consisting of H, alkyl, alkenyl, alkynyl, benzyl, and phenethyl;

or

A is

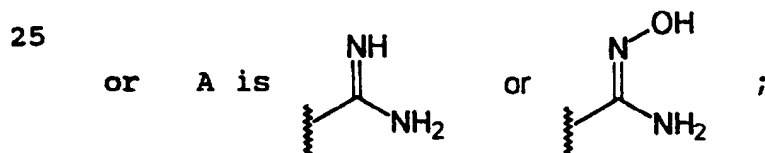


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wherein Y^2 is selected from the group consisting of H, alkyl; cycloalkyl; bicycloalkyl; aryl; monocyclic heterocycles; alkyl optionally substituted with aryl which can also be optionally substituted with one or more substituent selected from halo, haloalkyl, alkyl, nitro, hydroxy, alkoxy, aryloxy, aryl, or fused aryl; aryl optionally substituted with one or more substituent selected from halo, haloalkyl, hydroxy, alkoxy, aryloxy, aryl, fused aryl, nitro, methylenedioxy, ethylenedioxy, or alkyl; alkynyl; alkenyl; $-S-R^9$ and $-O-R^9$ wherein R^9 is selected from the group consisting of H; alkyl; aralkyl; aryl; alkenyl; and alkynyl; or R^9 taken together with R^7 forms a 4-12 membered mononitrogen containing sulfur or oxygen containing heterocyclic ring; and

R^5 and R^7 are as defined above;

or Y^2 (when Y^2 is carbon) taken together with R^7 forms a 4-12 membered mononitrogen containing ring optionally substituted with alkyl, aryl or hydroxy;

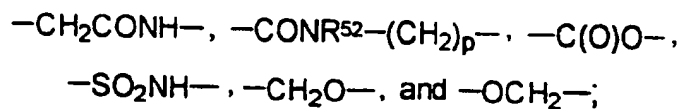


Z^1 , Z^2 , Z^4 and Z^5 are independently selected from the group consisting of H; alkyl; hydroxy; alkoxy; aryloxy; aralkoxy; halogen; haloalkyl; haloalkoxy; nitro; amino; aminoalkyl; alkylamino; dialkylamino; cyano; alkylthio; alkylsulfonyl; carboxyl derivatives; carboxyalkyl; alkoxy-carbonylalkyl; acetamide; aryl; fused aryl; cycloalkyl; thio; monocyclic heterocycles; fused

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monocyclic heterocycles; and A, wherein A is defined above;

5 B is selected from the group consisting of



10 wherein p is an integer selected from the group consisting of 0, 1 and 2;

wherein R^{50} is selected from the group consisting of H, alkyl, aryl, carboxyl derivatives and -
15 $\text{CONHCH}_2\text{CO}_2\text{R}^{53}$ wherein R^{53} is H or lower alkyl;

R^{52} is selected from the group consisting of H or alkyl;

20 R^{51} is selected from the group consisting of H, alkyl, carboxyl derivatives,

$$\begin{aligned} &-\text{NCO}_2\text{R}^{54}, -\text{NHSO}_2\text{R}^{54}, -\text{NHCONHR}^{54}, \\ &\text{H} \\ &-\text{NHCOR}^{54}, -\text{NHCOR}^{55} \text{ and amino;} \end{aligned}$$

25

wherein R^{54} is selected from the group consisting of H, alkyl, cycloalkyl, aryl, aralkyl, aralkenyl, and aryl substituted by one or more alkyl or halo;
30 wherein R^{55} is selected from the group consisting of N-substituted pyrrolidinyl, piperidinyl and morpholinyl;

t is an integer 0, 1 or 2;

35

R is X-R^3 wherein X is selected from the group consisting of O, S and NR^4 , wherein R^3 and R^4 are independently selected from the group consisting

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of hydrogen; alkyl; alkenyl; alkynyl; haloalkyl; aryl; arylalkyl; sugars; steroids and in the case of the free acid, all pharmaceutically acceptable salts thereof; and

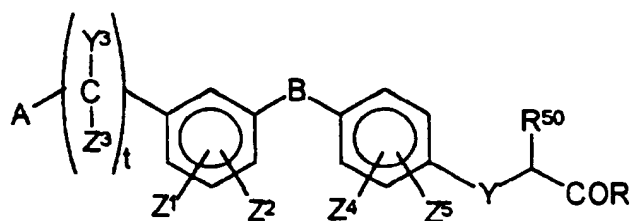
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Y^3 and Z^3 are independently selected from the group consisting of H, alkyl, aryl, cycloalkyl and aralkyl or Y^3 and Z^3 taken together with the C form a carbonyl.

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This invention also relates to a class of compounds represented by the Formula II

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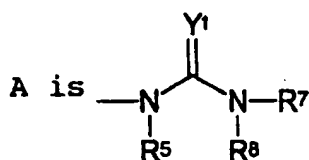


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or a pharmaceutically acceptable salt thereof, wherein

Y is selected from the group consisting of -O-, -S- and -SO₂-; wherein

25



30

wherein Y^1 is selected from the group consisting of N-R², O, and S;

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R^2 is selected from the group consisting of H; alkyl; aryl; hydroxy; alkoxy; cyano; nitro; amino; aminocarbonyl; alkenyl; alkynyl; alkyl optionally substituted with one or more substituent selected from lower alkyl, halogen, hydroxyl, haloalkyl, cyano, nitro, carboxyl, amino, alkoxy, aryl or aryl optionally substituted with one or more halogen, haloalkyl, lower alkyl, alkoxy, cyano, alkylsulfonyl, alkylthio, nitro, carboxyl, amino, hydroxyl, sulfonic acid, sulfonamide, aryl, fused aryl, monocyclic heterocycles, or fused monocyclic heterocycles; aryl optionally substituted with one or more substituent selected from halogen, haloalkyl, hydroxy, lower alkyl, alkoxy, methylenedioxy, ethylenedioxy, cyano, nitro, alkylthio, alkylsulfonyl, sulfonic acid, sulfonamide, carboxyl derivatives, amino, aryl, fused aryl, monocyclic heterocycles and fused monocyclic heterocycle; monocyclic heterocycles; and monocyclic heterocycles optionally substituted with one or more substituent selected from halogen, haloalkyl, lower alkyl, alkoxy, amino, nitro, hydroxy, carboxyl derivatives, cyano, alkylthio, alkylsulfonyl, sulfonic acid, sulfonamide, aryl or fused aryl; or

R^2 taken together with R^7 forms a 4-12 membered dinitrogen containing heterocycle optionally substituted with one or more substituent selected from the group consisting of lower alkyl, hydroxy and phenyl;

or R^2 taken together with R^7 forms a 5 membered heteroaromatic ring optionally substituted with amino;

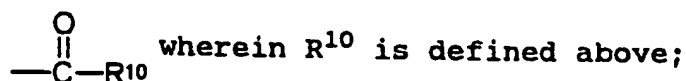
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or R^2 taken together with R^7 forms a 5 membered heteroaromatic ring optionally substituted with amino fused with a phenyl group;

- 5 R^7 (when not taken together with R^2) and R^8 are independently selected from the group consisting of H; alkyl; alkenyl; alkynyl; aralkyl; cycloalkyl; bicycloalkyl; aryl; acyl; benzoyl; alkyl optionally substituted with one or more
- 10 substituent selected from lower alkyl, halogen, hydroxy, haloalkyl, cyano, nitro, carboxyl derivatives, amino, alkoxy, thio, alkylthio, sulfonyl, aryl, aralkyl, aryl optionally substituted with one or more substituent selected
- 15 from halogen, haloalkyl, lower alkyl, alkoxy, methylenedioxy, ethylenedioxy, alkylthio, haloalkylthio, thio, hydroxy, cyano, nitro, carboxyl derivatives, aryloxy, amido, acylamino, amino, alkylamino, dialkylamino, trifluoroalkoxy, trifluoromethyl, sulfonyl, alkylsulfonyl,
- 20 haloalkylsulfonyl, sulfonic acid, sulfonamide, aryl, fused aryl, monocyclic heterocycles, fused monocyclic heterocycles; aryl optionally substituted with one or more substituent selected
- 25 from halogen, haloalkyl, lower alkyl, alkoxy, methylenedioxy, ethylenedioxy, alkylthio, haloalkylthio, thio, hydroxy, cyano, nitro, carboxyl derivatives, aryloxy, amido, acylamino, amino, alkylamino, dialkylamino, trifluoroalkoxy, trifluoromethylsulfonyl, alkylsulfonyl, sulfonic
- 30 acid, sulfonamide, aryl, fused aryl, monocyclic heterocycles, or fused monocyclic heterocycles; monocyclic heterocycles; monocyclic heterocycles optionally substituted with one or more
- 35 substituent selected from halogen, haloalkyl, lower alkyl, alkoxy, aryloxy, amino, nitro, hydroxy, carboxyl derivatives, cyano, alkylthio, alkylsulfonyl, aryl, fused aryl; monocyclic and

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bicyclic heterocyclicalkyls; $-\text{SO}_2\text{R}^{10}$ wherein R^{10} is selected from the group consisting of alkyl, aryl and monocyclic heterocycles, all optionally substituted with one or more substituent selected from the group consisting of halogen, haloalkyl, alkyl, alkoxy, cyano, nitro, amino, acylamino, trifluoroalkyl, amido, alkylaminosulfonyl, alkylsulfonyl, alkylsulfonylamino, alkylamino, dialkylamino, trifluoromethylthio, trifluoroalkoxy, trifluoromethylsulfonyl, aryl, aryloxy, thio, alkylthio, and monocyclic heterocycles; and

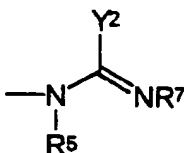


or NR^7 and R^8 taken together form a 4-12 membered mononitrogen containing monocyclic or bicyclic ring optionally substituted with one or more substituent selected from lower alkyl, carboxyl derivatives, aryl or hydroxy and wherein said ring optionally contains a heteroatom selected from the group consisting of O, N and S;

R^5 is selected from the group consisting of H, alkyl, alkenyl, alkynyl, benzyl, and phenethyl;

or

A is



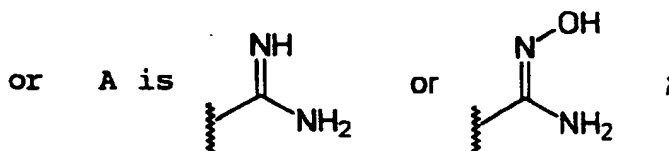
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wherein Y^2 is selected from the group consisting of alkyl; cycloalkyl; bicycloalkyl; aryl; monocyclic heterocycles; alkyl optionally substituted with aryl which can also be optionally substituted with one or more substituent selected from halo, haloalkyl, alkyl, nitro, hydroxy, alkoxy, aryloxy, aryl, or fused aryl; aryl optionally substituted with one or more substituent selected from halo, haloalkyl, hydroxy, alkoxy, aryloxy, aryl, fused aryl, nitro, methylenedioxy, ethylenedioxy, or alkyl; alkynyl; alkenyl; $-S-R^9$ and $-O-R^9$ wherein R^9 is selected from the group consisting of H; alkyl; aralkyl; aryl; alkenyl; and alkynyl; or R^9 taken together with R^7 forms a 4-12 membered mononitrogen containing sulfur or oxygen containing heterocyclic ring; and

R^5 and R^7 are as defined above;

or Y^2 (when Y^2 is carbon) taken together with R^7 forms a 4-12 membered mononitrogen containing ring optionally substituted with alkyl, aryl or hydroxy;

25



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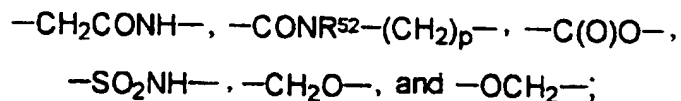
Z^1 , Z^2 , Z^4 and Z^5 are independently selected from the group consisting of H; alkyl; hydroxy; alkoxy; aryloxy; aralkoxy; halogen; haloalkyl; haloalkoxy; nitro; amino; aminoalkyl; alkylamino; dialkylamino; cyano; alkylthio; alkylsulfonyl; carboxyl derivatives; carboxyalkyl; alkoxycarbonylalkyl; acetamide; aryl; fused aryl;

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- 15 -

cycloalkyl; thio; monocyclic heterocycles; fused monocyclic heterocycles; and A, wherein A is defined above;

5 B is selected from the group consisting of



10 wherein p is an integer selected from the group consisting of 0, 1 and 2;

wherein R^{50} is selected from the group consisting of H, alkyl, aryl and carboxyl derivatives;

15 R^{52} is selected from the group consisting of H or alkyl;

t is an integer 0, 1 or 2;

20 R is X-R^3 wherein X is selected from the group consisting of O, S and NR^4 , wherein R^3 and R^4 are independently selected from the group consisting of hydrogen; alkyl; alkenyl; alkynyl; haloalkyl; aryl; arylalkyl; sugars; steroids and in the case
25 of the free acid, all pharmaceutically acceptable salts thereof; and

30 Y^3 and Z^3 are independently selected from the group consisting of H, alkyl, aryl, cycloalkyl and aralkyl or Y^3 and Z^3 taken together with C form a carbonyl.

35 It is another object of the invention to provide pharmaceutical compositions comprising compounds of the Formulas I and II. Such compounds and compositions are useful in selectively inhibiting or antagonizing the

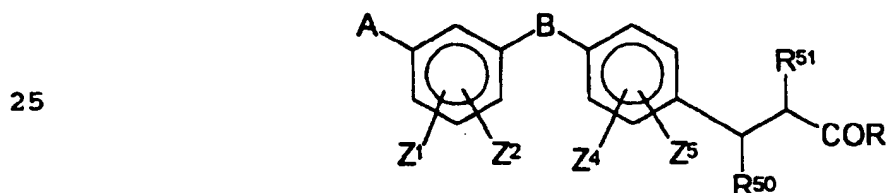
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$\alpha_v\beta_3$ and therefore in another embodiment the present invention relates to a method of selectively inhibiting or antagonizing the $\alpha_v\beta_3$ integrin. The invention further involves treating or inhibiting pathological conditions associated therewith such as osteoporosis, humoral hypercalcemia of malignancy, Paget's disease, tumor metastasis, solid tumor growth (neoplasia), angiogenesis, including tumor angiogenesis, retinopathy including diabetic retinopathy, arthritis, including rheumatoid arthritis, periodontal disease, psoriasis, smooth muscle cell migration and restenosis in a mammal in need of such treatment. Additionally, such pharmaceutical agents are useful as antiviral agents, and antimicrobials.

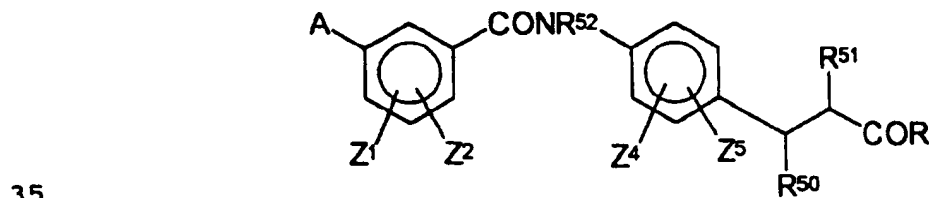
Detailed Description

The present invention relates to a class of compounds represented by the Formulas I and II, described above.

A preferred embodiment of the present invention is a compound of the Formula III

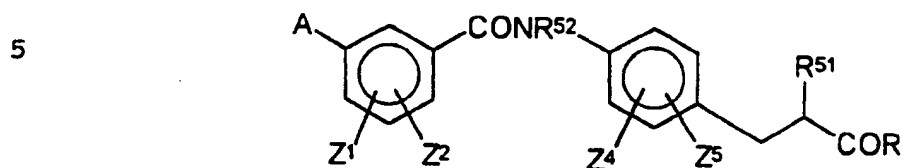


Another preferred embodiment of the present invention is a compound of the Formula IV

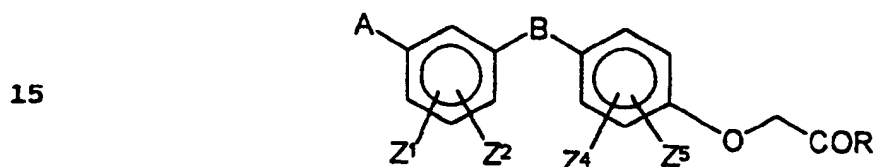


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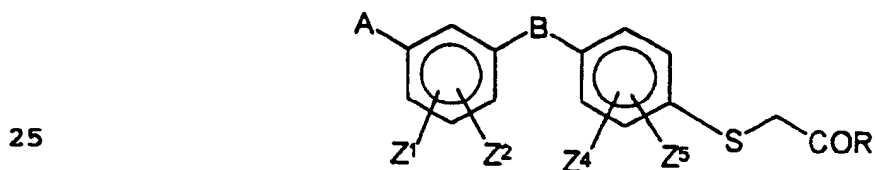
Another preferred embodiment of the present invention is a compound of the Formula V



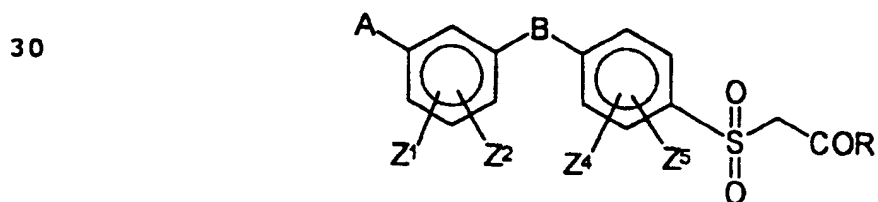
10 Another preferred embodiment of the present invention is a compound of the Formula VI



20 Another preferred embodiment of the present invention is a compound of the Formula VII

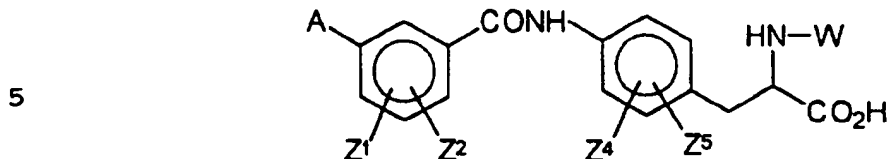


Another preferred embodiment of the present invention is a compound of the Formula VIII



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When the compound of the invention is of the formula



the compounds can be racemic or R or S stereochemistry. However, the S stereochemistry (derived from the natural amino acid, i.e. L-form) is the most preferred, followed by the racemic and followed by the R-stereoisomer.

The invention further relates to pharmaceutical compositions containing therapeutically effective amounts of the compounds of Formulas I-VIII.

The invention also relates to a method of selectively inhibiting or antagonizing the $\alpha_v\beta_3$ integrin and more specifically relates to a method of inhibiting bone resorption, periodontal disease, osteoporosis, humoral hypercalcemia of malignancy, Paget's disease, tumor metastasis, solid tumor growth (neoplasia), angiogenesis, including tumor angiogenesis, retinopathy including diabetic retinopathy, arthritis, including rheumatoid arthritis, smooth muscle cell migration and restenosis by administering a therapeutically effective amount of a compound of the Formula I-VIII to achieve such inhibition together with a pharmaceutically acceptable carrier.

The following is a list of definitions of various terms used herein:

As used herein, the terms "alkyl" or "lower alkyl" refer to a straight chain or branched chain hydrocarbon radicals having from about 1 to about 10 carbon atoms, and more preferably 1 to about 6 carbon atoms. Examples of such alkyl radicals are methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, sec-butyl, t-butyl, pentyl, neopentyl, hexyl, isohexyl, and the like.

As used herein the terms "alkenyl" or "lower alkenyl" refer to unsaturated acyclic hydrocarbon radicals containing at least one double bond and 2 to about 6 carbon atoms, which carbon-carbon double bond may have either cis or trans geometry within the alkenyl moiety, relative to groups substituted on the double bond carbons. Examples of such groups are ethenyl, propenyl, butenyl, isobutenyl, pentenyl, hexenyl and the like.

As used herein the terms "alkynyl" or "lower alkynyl" refer to acyclic hydrocarbon radicals containing one or more triple bonds and 2 to about 6 carbon atoms. Examples of such groups are ethynyl, propynyl, butynyl, pentynyl, hexynyl and the like.

The term "cycloalkyl" as used herein means saturated or partially unsaturated cyclic carbon radicals containing 3 to about 8 carbon atoms and more preferably 4 to about 6 carbon atoms. Examples of such cycloalkyl radicals include cyclopropyl, cyclopropenyl, cyclobutyl, cyclopentyl, cyclohexyl, 2-cyclohexen-1-yl, and the like.

The term "aryl" as used herein denotes aromatic ring systems composed of one or more aromatic rings. Preferred aryl groups are those consisting of one, two or three aromatic rings. The term embraces aromatic radicals such as phenyl, pyridyl, naphthyl, thiophene, furan, biphenyl and the like.

As used herein, the term "cyano" is represented by a radical of the formula $\text{—}\dot{\text{C}}\text{N}$.

The terms "hydroxy" and "hydroxyl" as used herein are synonymous and are represented by a radical of the formula $\text{—}\dot{\text{O}}\text{H}$.

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The term "lower alkylene" or "alkylene" as used herein refers to divalent linear or branched saturated hydrocarbon radicals of 1 to about 6 carbon atoms.

As used herein the term "alkoxy" refers to
5 straight or branched chain oxy containing radicals of the formula $-OR^{20}$, wherein R^{20} is an alkyl group as defined above. Examples of alkoxy groups encompassed include methoxy, ethoxy, n-propoxy, n-butoxy, isopropoxy, isobutoxy, sec-butoxy, t-butoxy and the
10 like.

As used herein the terms "arylalkyl" or "aralkyl" refer to a radical of the formula $\text{---}R^{22}-R^{21}$ wherein R^{21} is aryl as defined above and R^{22} is an alkylene as defined above. Examples of aralkyl groups include
15 benzyl, pyridylmethyl, naphthylpropyl, phenethyl and the like.

As used herein the term "aralkenyl" is represented by a radical of the formula $\text{---}R^{23}R^{21}$ where R^{21} is aryl as defined above and R^{23} is alkylene of 2-6 carbon atoms
20 with one or more C-C double bonds.

As used herein the term "aralkoxy" or "arylalkoxy" refers to a radical of the formula $\text{---}OR^{60}$ wherein R^{60} is aralkyl as defined above.

25 As used herein the term "nitro" is represented by a radical of the formula $\text{---}NO_2$.

As used herein the term "halo" or "halogen" refers to bromo, chloro, fluoro or iodo.

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As used herein the term "haloalkyl" refers to alkyl groups as defined above substituted with one or more of the same or different halo groups at one or more carbon atom. Examples of haloalkyl groups include
 5 trifluoromethyl, dichloroethyl, fluoropropyl and the like.

As used herein the term "N-substituted pyrrolidinyl, piperidinyl, and morpholinyl" refer respectively to radicals of the formula

10



As used herein the term "carboxyl" or "carboxy" refers to a radical of the formula -COOH.
 15

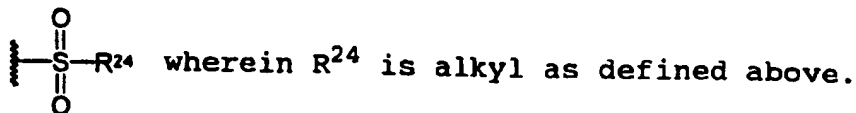
As used herein the term "carboxyl derivative" refers to a radical of the formula $\text{—}\overset{\text{Y}^6}{\underset{\text{||}}{\text{C}}}\text{—Y}^7\text{R}^{23}$ wherein

Y^6 and Y^7 are independently selected from the group consisting of O, N or S and R^{23} is selected from the group consisting of H, alkyl, aralkyl or aryl as
 20 defined above.

As used herein the term "amino" is represented by a radical of the formula -NH₂.

As used herein the term "aminoalkyl" refers to a
 25 radical of the formula $\text{H}_2\text{N—R}^{61}$ wherein R^{61} is alkylene as defined above.

As used herein the term "alkylsulfonyl" or "alkylsulfone" refers to a radical of the formula



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As used herein the term "alkylthio" refers to a radical of the formula $-SR^{24}$ wherein R^{24} is alkyl as defined above.

As used herein the term "sulfonic acid" refers to

5 a radical of the formula $\begin{array}{c} \text{O} \\ \parallel \\ \text{---S---OR}^{25} \\ \parallel \\ \text{O} \end{array}$ wherein R^{25} is H,

alkyl or aryl as defined above.

As used herein the term "sulfonamide" refers to a

radical of the formula $\begin{array}{c} \text{O} \\ \parallel \\ \text{---S---N} \begin{array}{l} \nearrow \text{R}^7 \\ \searrow \text{R}^8 \end{array} \\ \parallel \\ \text{O} \end{array}$ wherein R^7 and R^8 are as

defined above.

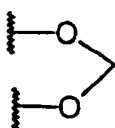
10 As used herein the term "fused aryl" refers to an aromatic ring such as the aryl groups defined above fused to one or more phenyl rings. Embraced by the term "fused aryl" is the radical naphthyl.

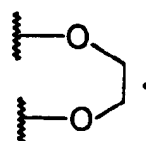
As used herein the terms "monocyclic heterocycle" or "monocyclic heterocyclic" refer to a monocyclic ring containing from 4 to about 12 atoms, and more preferably from 5 to about 10 atoms, wherein 1 to 3 of the atoms are heteroatoms selected from the group consisting of oxygen, nitrogen and sulfur with the understanding that if two or more different heteroatoms are present at least one of the heteroatoms must be nitrogen. Representative of such monocyclic heterocycles are imidazole, furan, pyridine, oxazole, pyran, triazole, thiophene, pyrazole, thiazole, thiadiazole, and the like.

As used herein the term "fused monocyclic heterocycle" refers to a monocyclic heterocycle as defined above with a benzene fused thereto. Examples of such fused monocyclic heterocycles include benzofuran, benzopyran, benzodioxole, benzothiazole, benzothiophene, benzimidazole and the like.

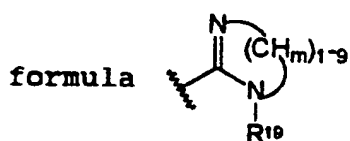
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As used herein the term "methylenedioxy" refers to

the radical  and the term "ethylenedioxy" refers

to the radical .

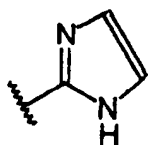
5 As used herein the term "4-12 membered dinitrogen containing heterocycle" refers to a radical of the



wherein m is 1 or 2 and R¹⁹ is

H, alkyl, aryl, or aralkyl and more preferably refers
to 4-9 membered ring and includes rings such as
10 imidazoline.

As used herein the term "5-membered heteroaromatic ring" includes for example a radical of the formula



and "5-membered heteroaromatic ring fused

with a phenyl" refers to such a "5-membered
15 heteroaromatic ring" with a phenyl fused thereto.
Representative of such 5-membered heteroaromatic rings
fused with a phenyl is benzimidazole.

As used herein the term "bicycloalkyl" refers to a
bicyclic hydrocarbon radical containing 6 to about 12
20 carbon atoms which is saturated or partially
unsaturated.

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As used herein the term "acyl" refers to a radical

of the formula $\begin{array}{c} \text{O} \\ \parallel \\ \text{---C---R}^{26} \end{array}$ wherein R^{26} is alkyl, alkenyl,

alkynyl, aryl or aralkyl as defined above. Encompassed
by such radical are the groups acetyl, benzoyl and the
5 like.

As used herein the term "thio" refers to a radical
of the formula $\begin{array}{c} \text{---S---} \\ | \\ | \\ | \end{array}$ SH.

As used herein the term "sulfonyl" refers to a

radical of the formula $\begin{array}{c} \text{O} \\ \parallel \\ \text{---S---R}^{27} \\ \parallel \\ \text{O} \end{array}$ wherein R^{27} is alkyl,

10 aryl or aralkyl as defined above.

As used herein the term "haloalkylthio" refers to
a radical of the formula ---S---R^{28} wherein R^{28} is haloalkyl
as defined above.

15 As used herein the term "aryloxy" refers to a
radical of the formula $\begin{array}{c} \text{---O---R}^{29} \\ | \\ | \\ | \end{array}$ wherein R^{29} is aryl as
defined above.

As used herein the term "acylamino" refers to a

radical of the formula $\begin{array}{c} \text{O} \\ \parallel \\ \text{R}^{30}\text{---C---NH---} \\ | \\ | \\ | \end{array}$ wherein R^{30} is alkyl,

20 aralkyl or aryl as defined above.

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As used herein the term "amido" refers to a radical of the formula $\text{---R}^{31}\text{---C(=O)NH}_2$ wherein R^{31} is a bond

or alkylene as defined above.

As used herein the term "alkylamino" refers to a radical of the formula ---NHR^{32} wherein R^{32} is alkyl as defined above.

As used herein the term "dialkylamino" refers to a radical of the formula $\text{---NR}^{33}\text{R}^{34}$ wherein R^{33} and R^{34} are the same or different alkyl groups as defined above.

As used herein the term "trifluoromethyl" refers to a radical of the formula ---CF_3 .

As used herein the term "trifluoroalkoxy" refers to a radical of the formula $\text{F}_3\text{C---R}^{35}\text{---O---}$ wherein R^{35} is a bond or an alkylene as defined above.

As used herein the term "alkylaminosulfonyl" refers to a radical of the formula $\text{R}^{36}\text{---N(H)---S(=O)}_2\text{---}$ wherein

R^{36} is alkyl as defined above.

As used herein the term "alkylsulfonylamino" refers to a radical of the formula $\text{R}^{36}\text{---S(=O)}_2\text{---NH---}$

wherein R^{36} is alkyl as defined above.

As used herein the term "trifluoromethylthio" refers to a radical of the formula $\text{F}_3\text{C---S---}$.

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As used herein the term "trifluoromethylsulfonyl"

refers to a radical of the formula $\text{F}_3\text{C}-\text{S}(=\text{O})_2-$.

As used herein the term "4-12 membered mono-nitrogen containing monocyclic or bicyclic ring" refers to a saturated or partially unsaturated monocyclic or bicyclic ring of 4-12 atoms and more preferably a ring of 4-9 atoms wherein one atom is nitrogen. Such rings may optionally contain additional heteroatoms selected from nitrogen, oxygen or sulfur. Included within this group are morpholine, piperidine, piperazine, thiomorpholine, pyrrolidine, proline, azacycloheptene and the like.

As used herein the term "benzyl" refers to the

radical $-\text{CH}_2-\text{C}_6\text{H}_5$.

As used herein the term "phenethyl" refers to the

radical $-\text{CH}_2\text{CH}_2-\text{C}_6\text{H}_5$.

As used herein the term "4-12 membered mono-nitrogen containing sulfur or oxygen containing heterocyclic ring" refers to a ring consisting of 4 to 12 atoms and more preferably 4 to 9 atoms wherein at least one atom is a nitrogen and at least one atom is oxygen or sulfur. Encompassed within this definition are rings such as thiazoline and the like.

As used herein the term "arylsulfonyl" or "arylsulfone" refers to a radical of the formula

$\text{R}^{37}-\text{S}(=\text{O})_2-$ wherein R^{37} is aryl as defined above.

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As used herein the terms "alkylsulfoxide" or "arylsulfoxide" refer to radicals of the formula



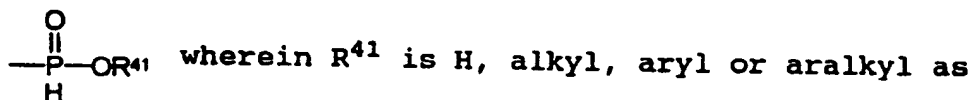
defined above.

5 As used herein the term "phosphonic acid

derivative" refers to a radical of the formula $\text{[radical]}-\overset{\text{O}}{\parallel}{\text{P}}(\text{OR}^{39})(\text{OR}^{40})$

wherein R^{39} and R^{40} are the same or different H, alkyl, aryl or aralkyl.

10 As used herein the term "phosphinic acid derivatives" refers to a radical of the formula



defined above.

As used herein the term "arylthio" refers to a radical of the formula $\text{[radical]}-\text{SR}^{42}$ wherein R^{42} is aryl as

15 defined above.

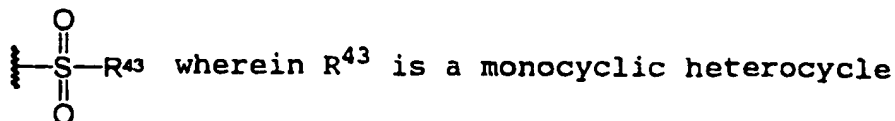
As used herein the term "monocyclic heterocycle thio" refers to a radical of the formula $\text{[radical]}-\text{SR}^{43}$

wherein R^{43} is a monocyclic heterocycle radical as defined above.

20 As used herein the terms "monocyclic heterocycle sulfoxide" and "monocyclic heterocycle sulfone" refer,

respectively, to radicals of the formula $\text{[radical]}-\overset{\text{O}}{\parallel}{\text{S}}-\text{R}^{43}$ and

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radical as defined above.

The term "composition" as used herein means a product which results from the mixing or combining of more than one element or ingredient.

The term "pharmaceutically acceptable carrier", as used herein means a pharmaceutically-acceptable material, composition or vehicle, such as a liquid or solid filler, diluent, excipient, solvent or encapsulating material, involved in carrying or transporting a chemical agent.

The term "therapeutically effective amount" shall mean that amount of drug or pharmaceutical agent that will elicit the biological or medical response of a tissue, system or animal that is being sought by a researcher or clinician.

The following is a list of abbreviations and the corresponding meanings as used interchangeably herein:

¹H-NMR = proton nuclear magnetic resonance
 AcOH = acetic acid
 BH₃-THF = borane-tetrahydrofuran complex
 BOC = tert-butoxycarbonyl
 Cat. = catalytic amount
 CH₂Cl₂ = dichloromethane
 CH₃CN = acetonitrile
 CH₃I = iodomethane
 CHN analysis = carbon/hydrogen/nitrogen elemental analysis
 CHNCl analysis = carbon/hydrogen/nitrogen/chlorine elemental analysis
 CHNS analysis = carbon/hydrogen/nitrogen/sulfur elemental analysis
 DCC = 1,3-dicyclohexylcarbodiimide
 DIEA = diisopropylethylamine
 DMA = N,N-dimethylacetamide
 DMAP = 4-(N,N-dimethylamino)pyridine
 DMF = N,N-dimethylformamide
 DSC = disuccinyl carbonate

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EDCI = 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride
Et₂O = diethyl ether
Et₃N = triethylamine
5 EtOAc = ethyl acetate
EtOH = ethanol
FAB MS = fast atom bombardment mass spectroscopy
g = gram(s)
10 GIHA HCl = meta-guanidino-hippuric acid hydrochloride
GIHA = meta-guanidino-hippuric acid
HPLC = high performance liquid chromatography
IBCF = isobutylchloroformate
15 K₂CO₃ = potassium carbonate
KOH = potassium hydroxide
LiOH = lithium hydroxide
MCPBA = m-chloroperoxybenzoic acid or m-chloroperbenzoic acid
20 MeOH = methanol
MesCl = methanesulfonylchloride
mg = milligram
MgSO₄ = magnesium sulfate
ml = milliliter
mL = milliliter
25 MS = mass spectroscopy
N₂ = nitrogen
NaCNBH₃ = sodium cyanoborohydride
Na₂PO₄ = sodium phosphate
Na₂SO₄ = sodium sulfate
30 NaHCO₃ = sodium bicarbonate
NaOH = sodium hydroxide
NH₄HCO₃ = ammonium bicarbonate
NH₄⁺HCO₂⁻ = ammonium formate
35 NMM = N-methyl morpholine
NMR = nuclear magnetic resonance
RPHPLC = reverse phase high performance liquid chromatography
RT = room temperature
KSCN = potassium thiocyanate
40 Pd/C = palladium on carbon
Bn = benzyl
Et = ethyl
Me = methyl
Ph = phenyl
45 NEt₃ = triethylamine
t-BOC = tert-butoxycarbonyl
TFA = trifluoroacetic acid
THF = tetrahydrofuran
50 Δ = heating the reaction mixture

As used herein HPLC-Method 1 refers to reverse phase C-18 functionalized silica gel column (50 x 300 mm) using a linear gradient of 95% 0.6% TFA/water:5%

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CH₃CN to 60% 0.6% TFA/water: 40% CH₃CN with a flow rate of 80 ml/minute.

The compounds as shown in Formulas I-VIII can exist in various isomeric forms and all such isomeric forms are meant to be included. Tautomeric forms are also included as well as pharmaceutically acceptable salts of such isomers and tautomers.

In the structures and formulas herein, a bond drawn across a bond of a ring can be to any available atom on the ring.

The term "pharmaceutically acceptable salt" refers to a salt prepared by contacting a compound of Formula I or II with an acid whose anion is generally considered suitable for human consumption. Examples of pharmacologically acceptable salts include the hydrochloride, hydrobromide, hydroiodide, sulfate, phosphate, acetate, propionate, lactate, maleate, malate, succinate, tartrate salts and the like. All of the pharmacologically acceptable salts may be prepared by conventional means. (See Berge et al., J Pharm. Sci., 66(1), 1-19 (1977) for additional examples of pharmaceutically acceptable salts.)

For the selective inhibition or antagonism of $\alpha_v\beta_3$ integrins, compounds of the present invention may be administered orally, parenterally, or by inhalation spray, or topically in unit dosage formulations containing conventional pharmaceutically acceptable carriers, adjuvants and vehicles. The term parenteral as used herein includes, for example, subcutaneous, intravenous, intramuscular, intrasternal, infusion techniques or intraperitoneally.

The compounds of the present invention are administered by any suitable route in the form of a pharmaceutical composition adapted to such a route, and in a dose effective for the treatment intended. Therapeutically effective doses of the compounds required to prevent or arrest the progress of or to

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treat the medical condition are readily ascertained by one of ordinary skill in the art using preclinical and clinical approaches familiar to the medicinal arts.

Accordingly, the present invention provides a
5 method of treating conditions mediated by selectively inhibiting or antagonizing the $\alpha_v\beta_3$ cell surface receptor which method comprises administering a therapeutically effective amount of a compound selected from the class of compounds depicted in Formulas I-
10 VIII, wherein one or more compounds of the Formulas I-VIII is administered in association with one or more non-toxic, pharmaceutically acceptable carriers and/or diluents and/or adjuvants (collectively referred to herein as "carrier" materials) and if desired other
15 active ingredients. More specifically, the present invention provides a method for inhibition of the $\alpha_v\beta_3$ cell surface receptor. Most preferably the present invention provides a method for inhibiting bone resorption, treating osteoporosis, inhibiting humoral
20 hypercalcemia of malignancy, treating Paget's disease, inhibiting tumor metastasis, inhibiting neoplasia (solid tumor growth), inhibiting angiogenesis including tumor angiogenesis, treating diabetic retinopathy, inhibiting arthritis, psoriasis and periodontal
25 disease, and inhibiting smooth muscle cell migration including restenosis.

Based upon standard laboratory experimental techniques and procedures well known and appreciated by those skilled in the art, as well as comparisons with
30 compounds of known usefulness, the compounds of Formula I can be used in the treatment of patients suffering from the above pathological conditions. One skilled in the art will recognize that selection of the most appropriate compound of the invention is within the
35 ability of one with ordinary skill in the art and will depend on a variety of factors including assessment of results obtained in standard assay and animal models.

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Treatment of a patient afflicted with one of the pathological conditions comprises administering to such a patient an amount of compound of the Formula I or II which is therapeutically effective in controlling the condition or in prolonging the survivability of the patient beyond that expected in the absence of such treatment. As used herein, the term "inhibition" of the condition refers to slowing, interrupting, arresting or stopping the condition and does not necessarily indicate a total elimination of the condition. It is believed that prolonging the survivability of a patient, beyond being a significant advantageous effect in and of itself, also indicates that the condition is beneficially controlled to some extent.

As stated previously, the compounds of the invention can be used in a variety of biological, prophylactic or therapeutic areas. It is contemplated that these compounds are useful in prevention or treatment of any disease state or condition wherein the $\alpha_v\beta_3$ integrin plays a role.

The dosage regimen for the compounds and/or compositions containing the compounds is based on a variety of factors, including the type, age, weight, sex and medical condition of the patient; the severity of the condition; the route of administration; and the activity of the particular compound employed. Thus the dosage regimen may vary widely. Dosage levels of the order from about 0.01 mg to about 1000 mg per kilogram of body weight per day are useful in the treatment of the above-indicated conditions and more preferably of the order from about 0.01 mg to about 100 mg per kilogram.

The active ingredient administered by injection is formulated as a composition wherein, for example, saline, dextrose or water may be used as a suitable carrier. A suitable daily dose would typically be

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about 0.01 to 100 mg/kg body weight injected per day in multiple doses depending on the factors listed above and more preferably of the order from about 0.01 to about 10 mg/kg.

5 For administration to a mammal in need of such treatment, the compounds in a therapeutically effective amount are ordinarily combined with one or more adjuvants appropriate to the indicated route of administration. The compounds may be admixed with
10 lactose, sucrose, starch powder, cellulose esters of alkanolic acids, cellulose alkyl esters, talc, stearic acid, magnesium stearate, magnesium oxide, sodium and calcium salts of phosphoric and sulphuric acids, gelatin, acacia, sodium alginate, polyvinylpyrrolidone,
15 and/or polyvinyl alcohol, and tableted or encapsulated for convenient administration. Alternatively, the compounds may be dissolved in water, polyethylene glycol, propylene glycol, ethanol, corn oil, cottonseed oil, peanut oil, sesame oil, benzyl alcohol, sodium
20 chloride, and/or various buffers. Other adjuvants and modes of administration are well and widely known in the pharmaceutical art.

 The pharmaceutical compositions useful in the present invention may be subjected to conventional
25 pharmaceutical operations such as sterilization and/or may contain conventional pharmaceutical adjuvants such as preservatives, stabilizers, wetting agents, emulsifiers, buffers, etc.

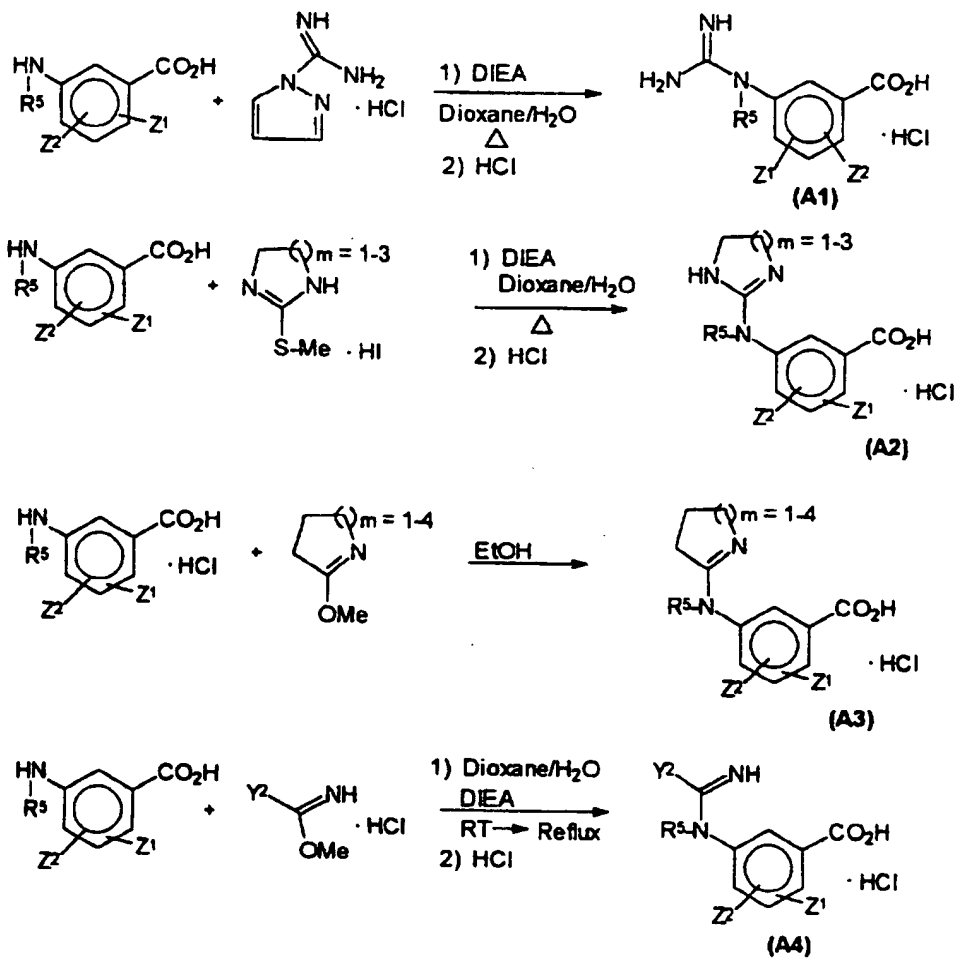
 The general synthetic sequences for preparing the
30 compounds useful in the present invention are outlined in Schemes I-IV. Both an explanation of, and the actual procedures for, the various aspects of the present invention are described where appropriate. The following Schemes and Examples are intended to be
35 merely illustrative of the present invention, and not limiting thereof in either scope or spirit. Those of skill in the art will readily understand that known

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variations of the conditions and processes described in the Schemes and Examples can be used to perform the process of the present invention.

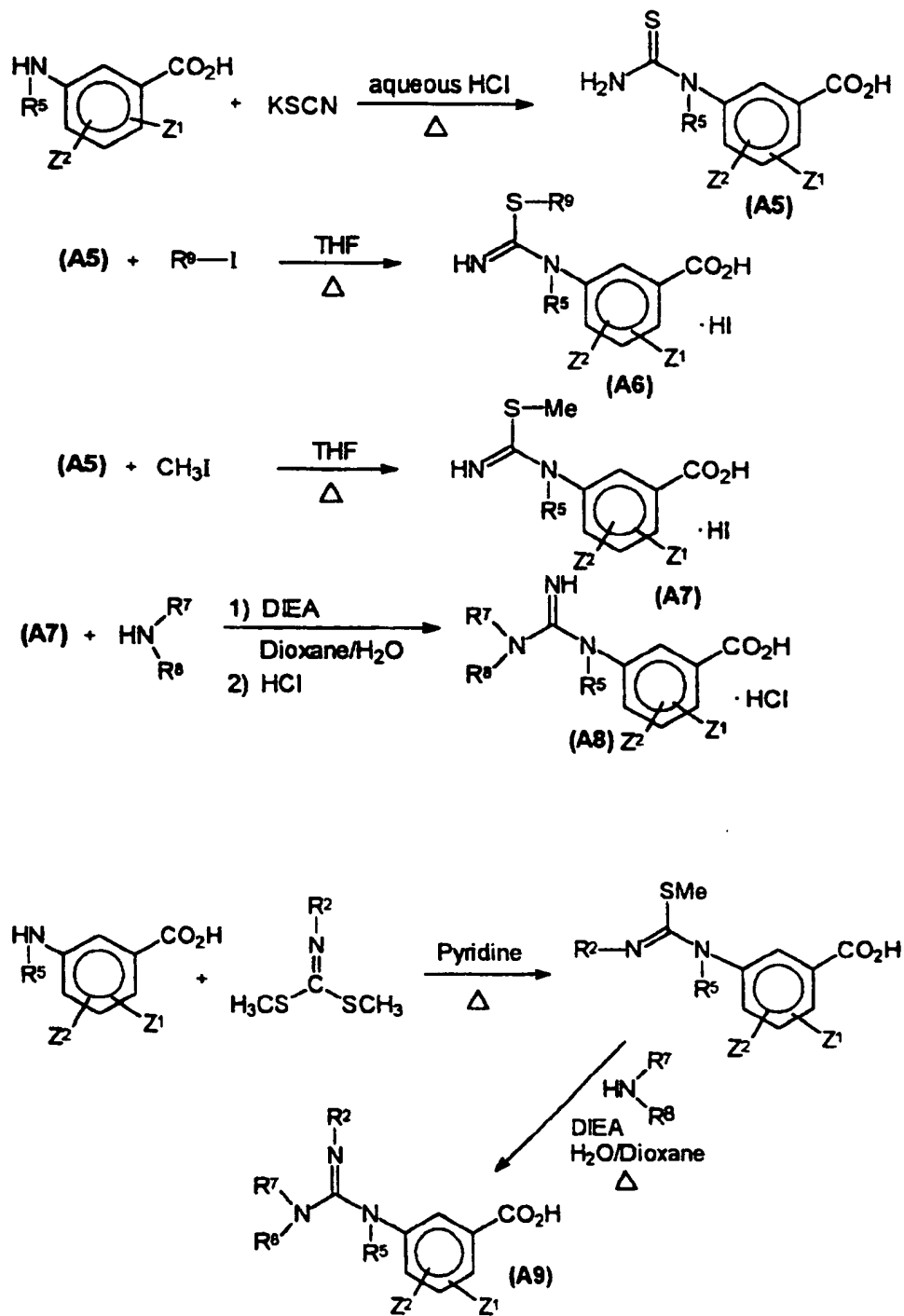
Unless otherwise indicated all starting materials
5 and equipment employed were commercially available.

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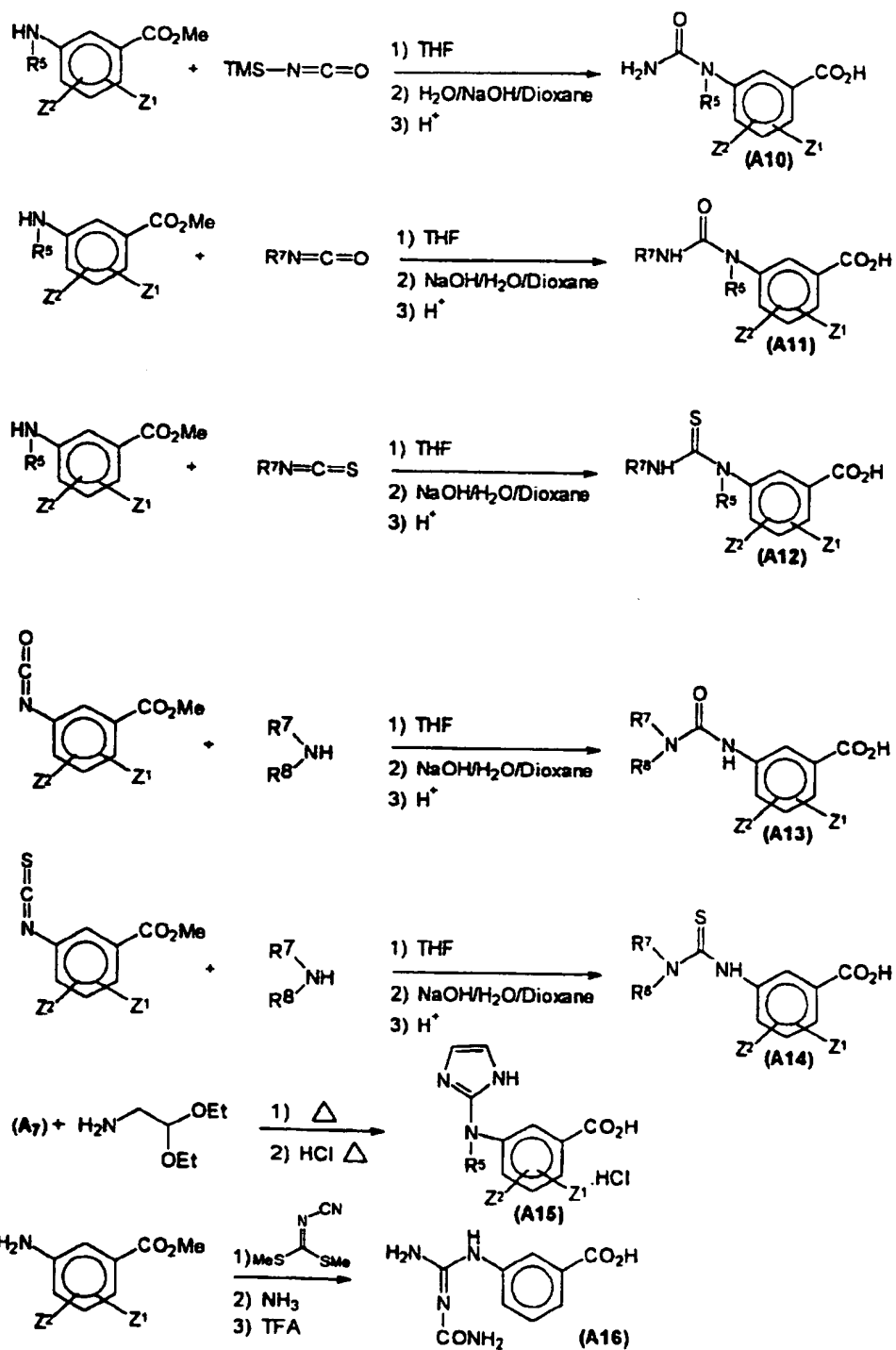
SCHEME I

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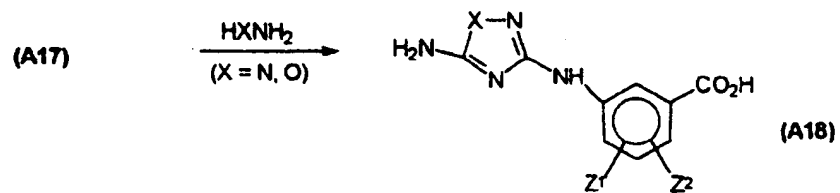
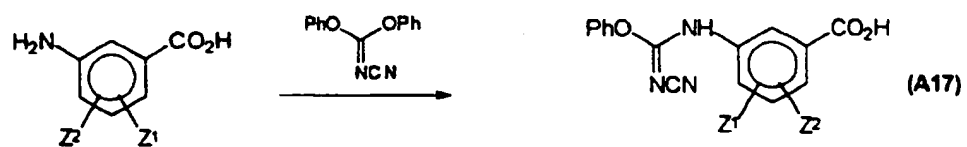
SCHEME I (Cont'd)



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SCHEME I (Cont'd)

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SCHEME I (Cont'd)

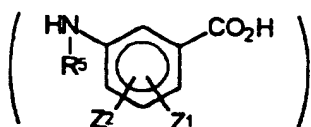
- 39 -

Schemes I, II and III are illustrative of methodology useful for preparing various compounds of the present invention. Such methodology is more specifically defined in the examples which follow.

5 Such methodology can be modified by one skilled in the art, substituting known reagents and conditions from conventional methodology to produce the desired compounds.

Specifically, in Scheme I:

10 In the synthesis of intermediate benzoic acids (A1) through (A16), the starting amino benzoic acids



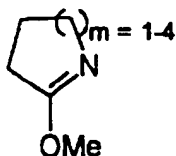
are either commercially available or

15 can be converted to such amino benzoic acids via reduction of the corresponding nitro benzoic acid, which can be obtained commercially or synthesized by nitration of the appropriate benzoic acid, followed by reduction to the desired amino benzoic acid. These are all when R⁵ is H. If R⁵ is other than H, alkylation of
20 the amino functionality can be achieved by conventional methodology.

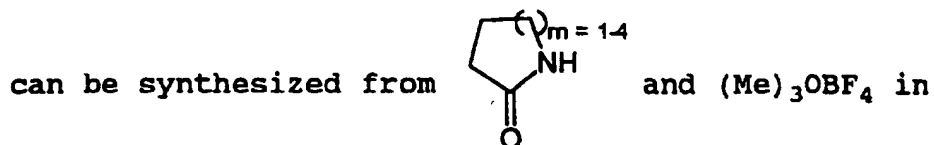
Furthermore, synthesis of intermediate (A2) can also be accomplished as disclosed generally in US 3,202,660, starting with the appropriate amino benzoic
25 acid. Furthermore, intermediate (A2) and (A15) as well as further analogues of (A2) and (A15) such as substitutions on the heterocyclic ring, oxazolidines, thiazolidines, benzimidazoles and the like can also be accomplished as disclosed in

- 30
- 1) Chem. Pharm. Bull. 41(1) 117-125 (1993)
 - 2) Chem. Pharm. Bull. 33(10) 4409-4421 (1985)
 - 3) J. Med. Chem. 18 (1), 90-99 (1975).

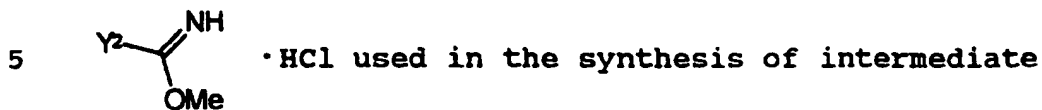
- 40 -



used in the synthesis of intermediates (A3),



dichloromethane.

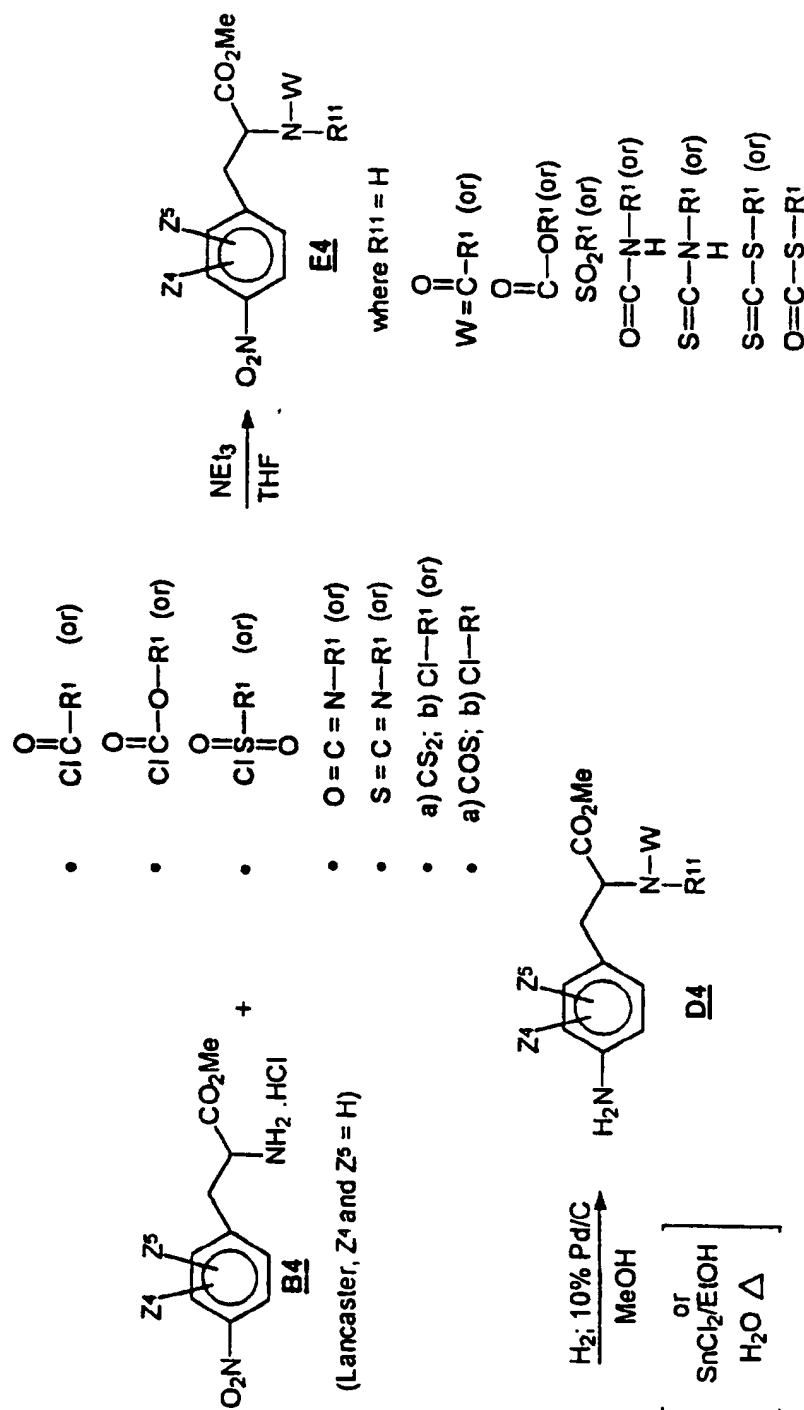


(A4), can be synthesized from Y²-CN and MeOH (1 equivalent) and HCl gas (1 equivalent) in heptane.

All other reagents in Scheme I are either
commercially available or readily synthesized by
10 methodologies known by those skilled in the art.

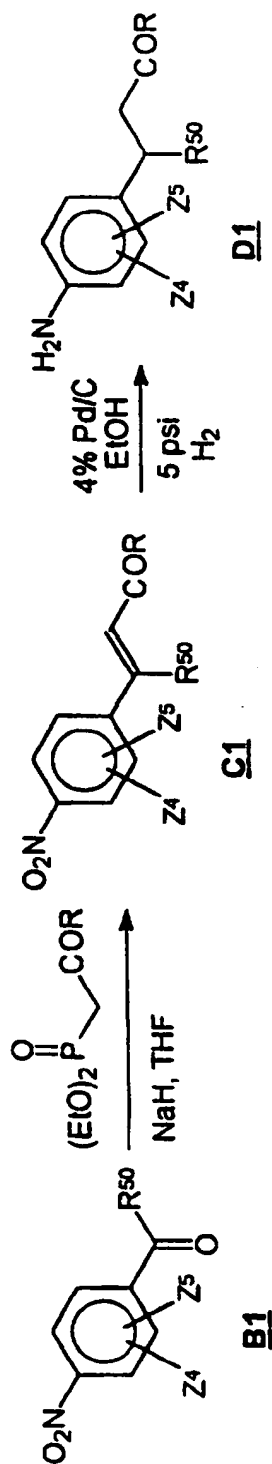
SCHEME II

(A)

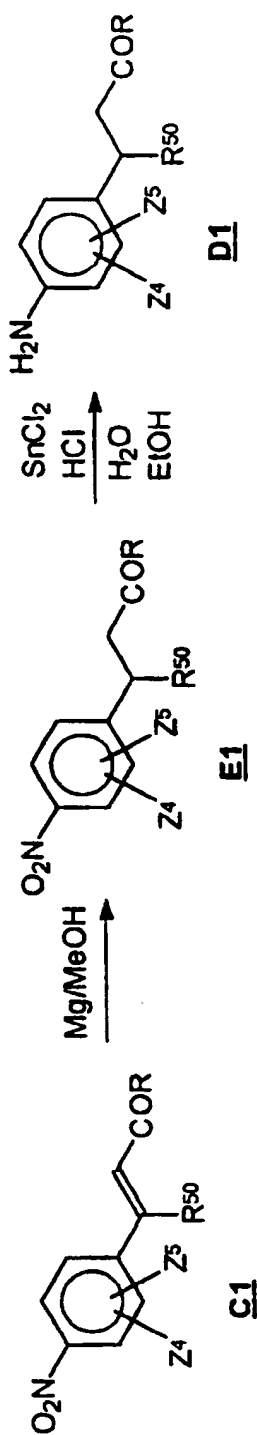


SCHEME II (cont'd)

(B)



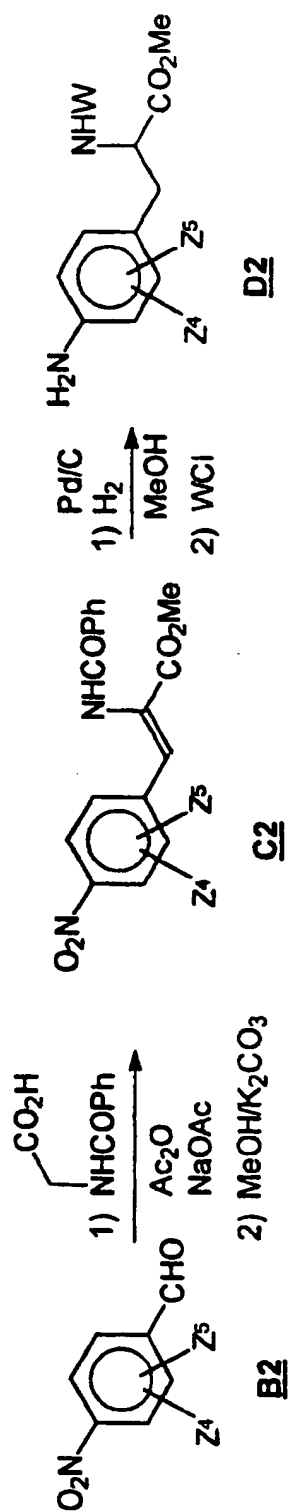
(C)



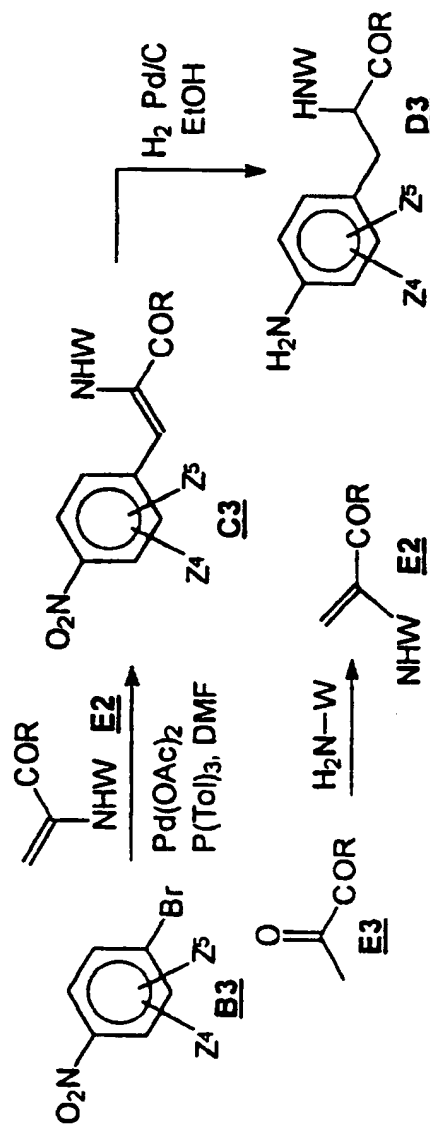
- 43 -

SCHEME II (cont'd)

(D)



(E)



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In Scheme II(A), phenylpropionic acid analogs D4 are readily prepared from 4-nitrophenyl alanine methyl esters B4 using the following procedure.

Nitrophenyl alanine B4 condenses with a variety of electrophiles such as R^1COCl , R^1OCOCl , R^1SO_2Cl , $R^1N=C=O$, $R^1N=C=S$, R^1SCOCl under standard conditions (NEt_3 , THF) known in the art. The resulting intermediate E4 is reduced by either catalytic hydrogenation (H_2 , Pd/C, MeOH), or tin II chloride ($SnCl_2$, H_2O , HCl, EtOH 100°) to afford the desired phenylpropionic acid analog D4.

In Scheme II(B) phenylpropionic acid analog D1 is readily prepared from aldehyde/or ketone B1 in the following manner.

Aldehyde or ketone B1 is condensed with $(EtO)_2P(O)CH_2COR$ under standard conditions (NaH/THF 0° to room temperature). The resulting cinnamic acid derivative C1 is reduced (4% Pd/C, EtOH, 5 psi) to afford the desired phenylpropionic acid analogs D1.

When substituents Z^4 and Z^5 are sensitive to the catalytic hydrogenation conditions described above, the following synthetic procedure depicted in Scheme II(C) may be utilized.

Nitrophenylcinnamic acid derivative C1 is partially reduced with magnesium in MeOH to afford nitrophenylpropionic acid analog E1, further reduction of the nitro moiety ($SnCl_2/H_2O/HCl/EtOH$) affords the desired phenylpropionic acid derivative D1.

In Scheme II(D) phenylpropionic acid derivative D2 can be readily prepared from aldehyde B2 as described below.

Aldehyde B2 is condensed with N-benzoyl glycine ($Ac_2O/100^\circ$) and the resulting azalactone is hydrolysed ($MeOH/K_2CO_3$) to afford the corresponding dehydroamino acid analog C2. Hydrogenation of C2 (Pd/C, H_2 , MeOH, 60 psi) followed by derivatisation with the electrophilic reagents W-Cl (as described in Scheme II(A)) affords phenylpropionic acid derivative D2.

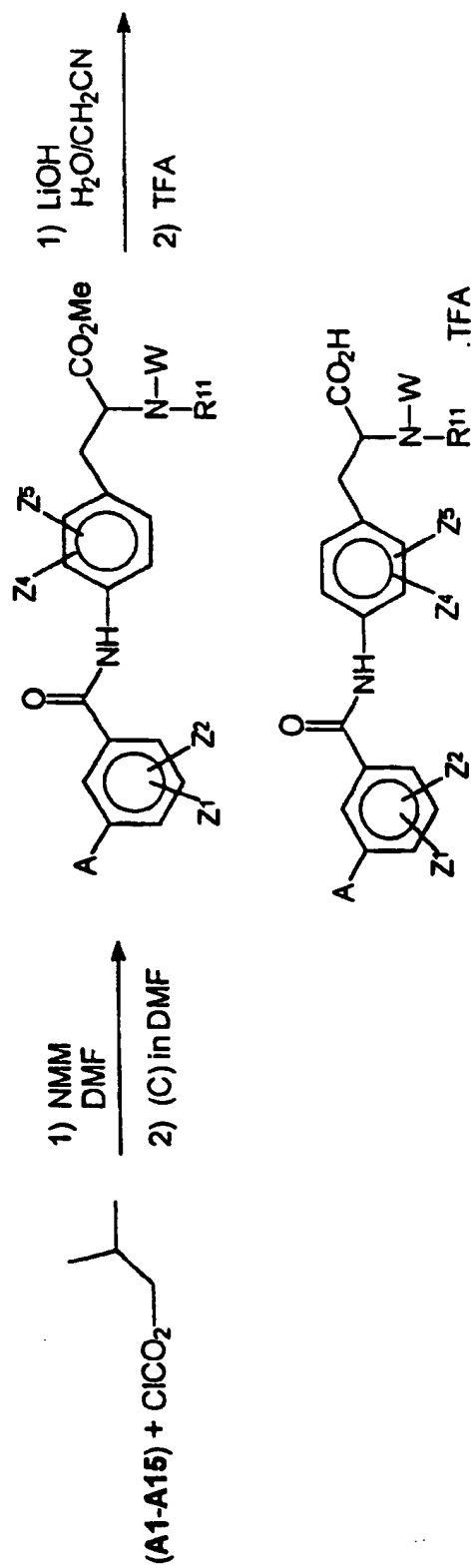
- 45 -

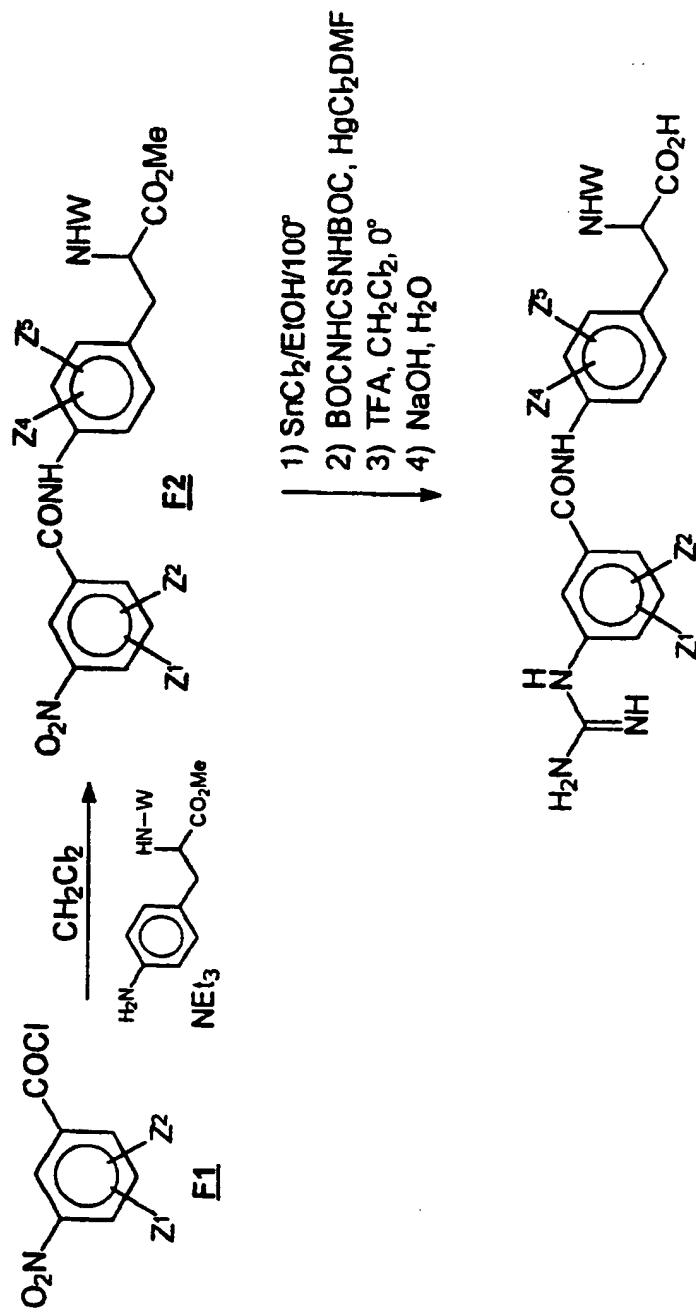
In Scheme II(E), phenylpropionic acid analog D3 may be prepared from bromide B3.

Bromide B3 is readily coupled to acrylate E2 using standard Heck coupling procedures such as ($\text{Pd}(\text{OAc})_2$, $\text{P}(\text{Tol})_3$, DMF, 130°) to afford the corresponding dehydroamino acid derivative C3. Hydrogenation of C3 (Pd/C , H_2 , EtOH, 60 psi) affords the desired phenylpropionic acid analog B3.

Furthermore acrylates E2 are readily prepared from condensation of pyruvic acid derivatives with $\text{H}_2\text{N-W}$ using standard dehydrating conditions such as (Tol, POCl_3).

Coupling of the intermediates from Scheme I [(A1) through (A16)] with the intermediates (D1-D4) (from Scheme II) can be accomplished using other coupling reagents known to those in the art in addition to the mixed anhydride method described in Scheme III, to give the final desired products.

SCHEME III**Method A**

SCHEME III (Cont'd)Method B

wherein W is as defined in Scheme II

In Scheme III - Method B

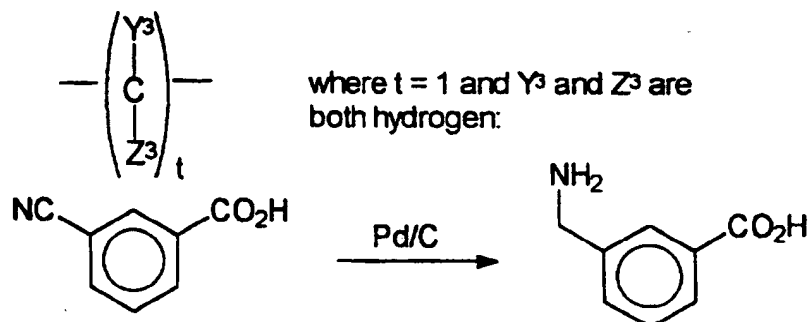
An alternative method to prepare compounds of the present invention is described below.

In this procedure, intermediates **D1-D4** [from Scheme II (A-F)] are coupled to 3-nitrobenzoylchloride **F1** (CH_2Cl_2 , NEt_3 , 0°). The resulting coupled product **F2** is reduced ($\text{SnCl}_2/\text{EtOH}$, H_2O , 100°) to the corresponding aniline. The resulting aniline can be converted into compounds of the present invention using the reactions described in Scheme I (A1-A6) or reacted with protected cyclic or acyclic thioureas, followed by deprotection ($\text{TFA}/\text{CH}_2\text{Cl}_2/0^\circ$).

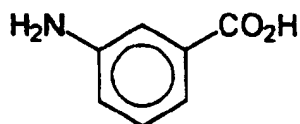
This procedure is exemplified by converting the above aniline to its correspondence guanidine analog (BOCNHCSNHBOC , HgCl_2 , DMF) followed by deprotection (TFA , CH_2Cl_2).

When R^{11} is not H, the appropriate nitrogen can be alkylated in an appropriate step by methodology known to those skilled in the art. Alternate acid derivatives R are synthesized by methodologies known to those skilled in the art.

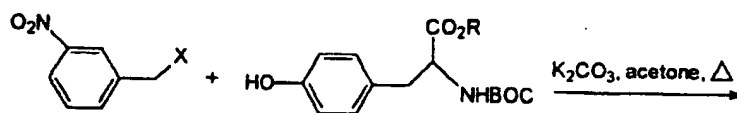
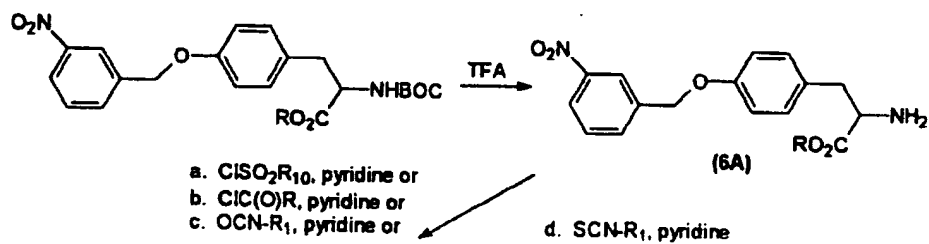
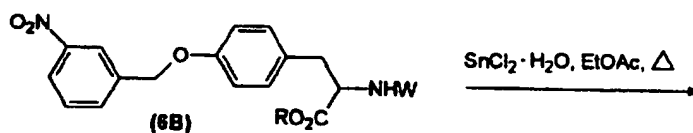
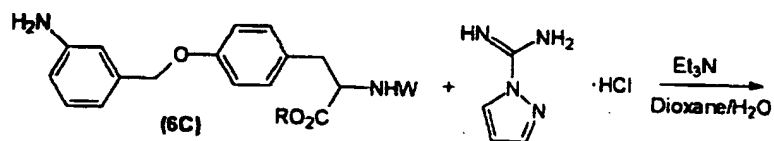
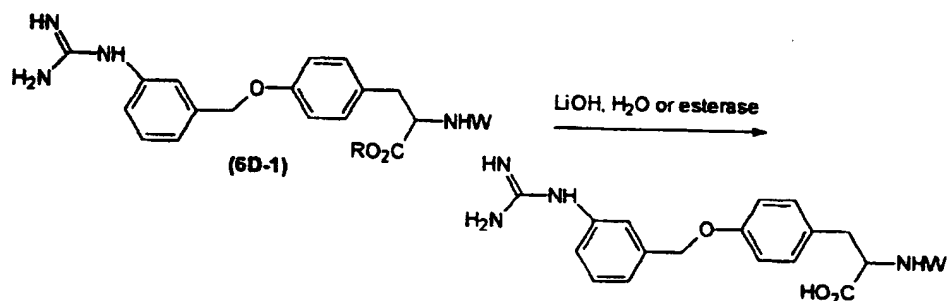
To synthesize compounds wherein



which is then treated in the same manner of further derivatization as exemplified in the previous schemes for:



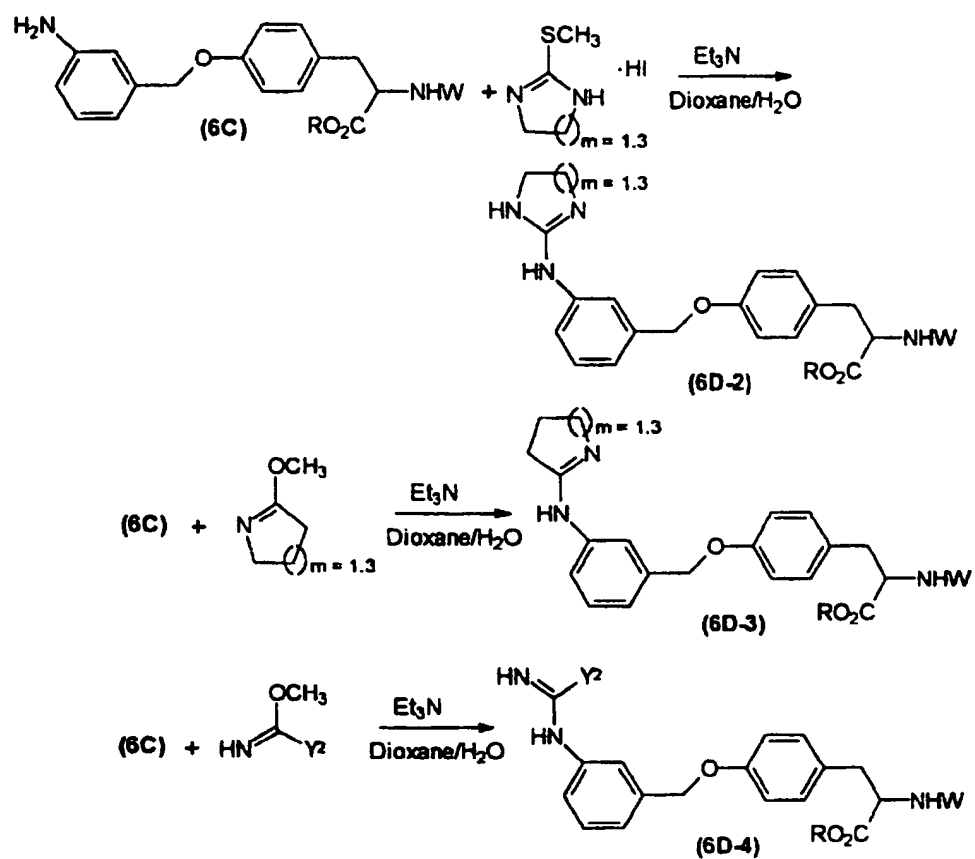
- 49 -

SCHEME IVSTEP 1 $\text{X} = \text{Cl, Br, OSO}_2\text{R}$ STEP 2STEP 3STEP 4STEP 5

- 50 -

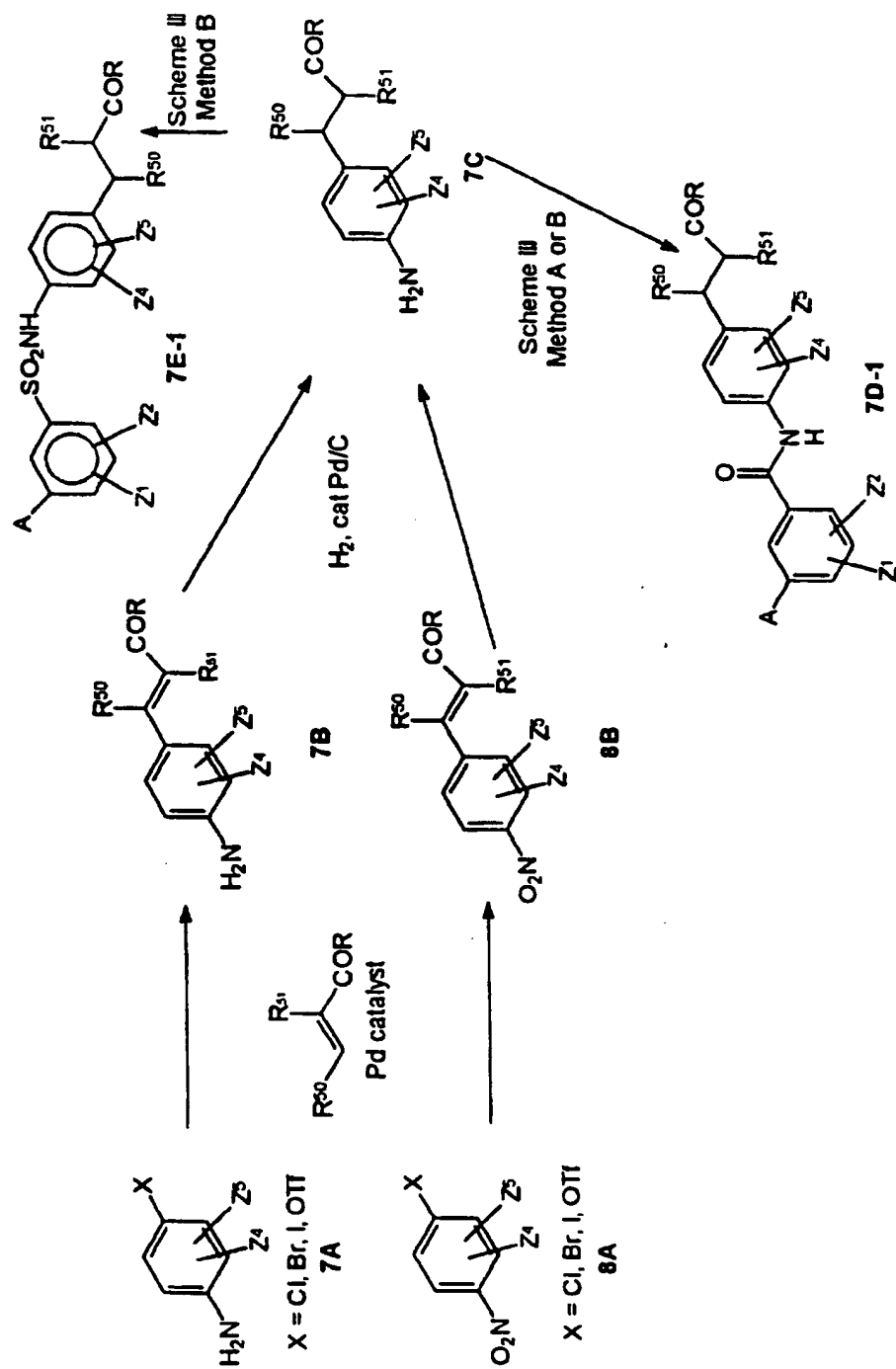
SCHEME IV (Cont'd)

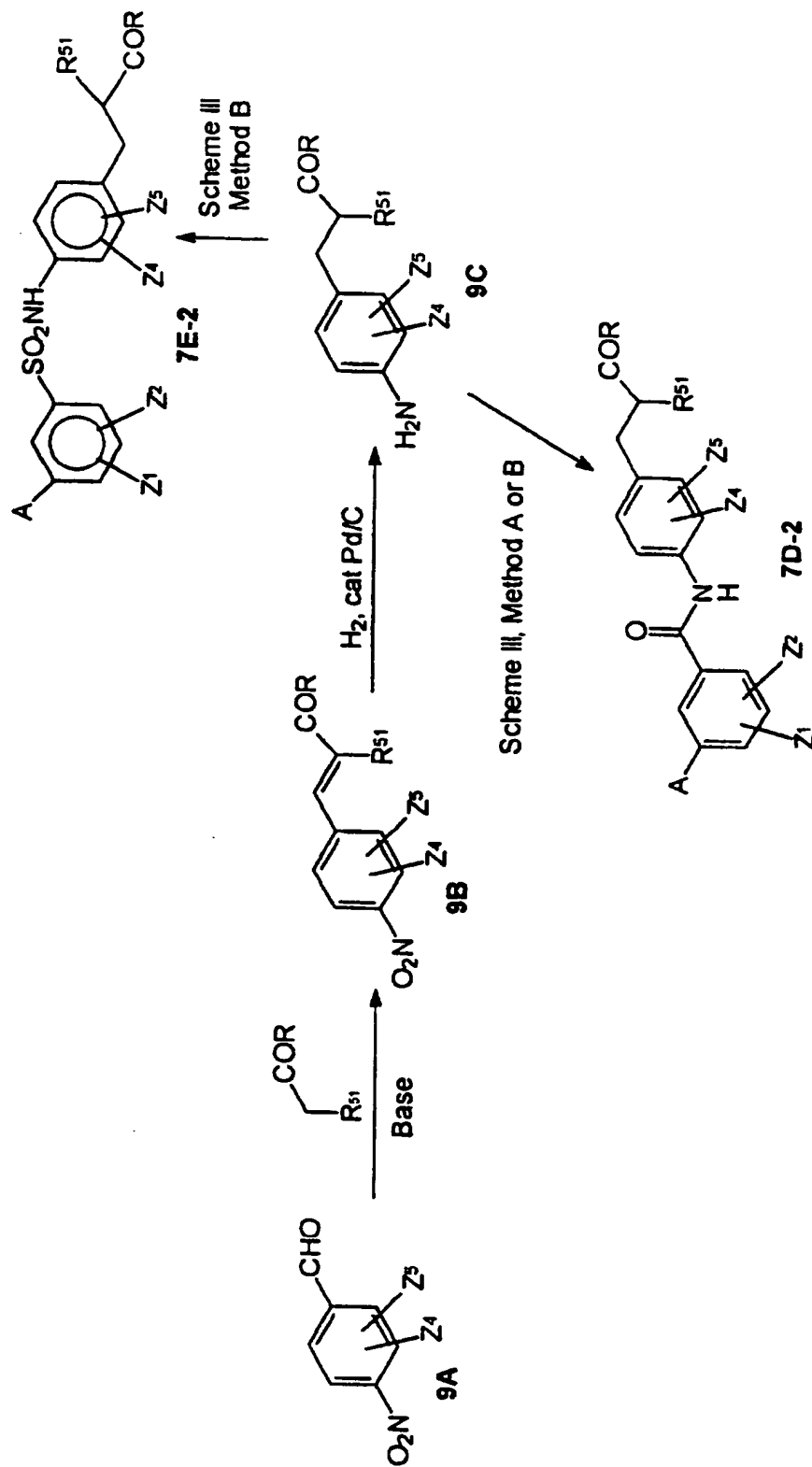
STEP 4 (Cont'd)



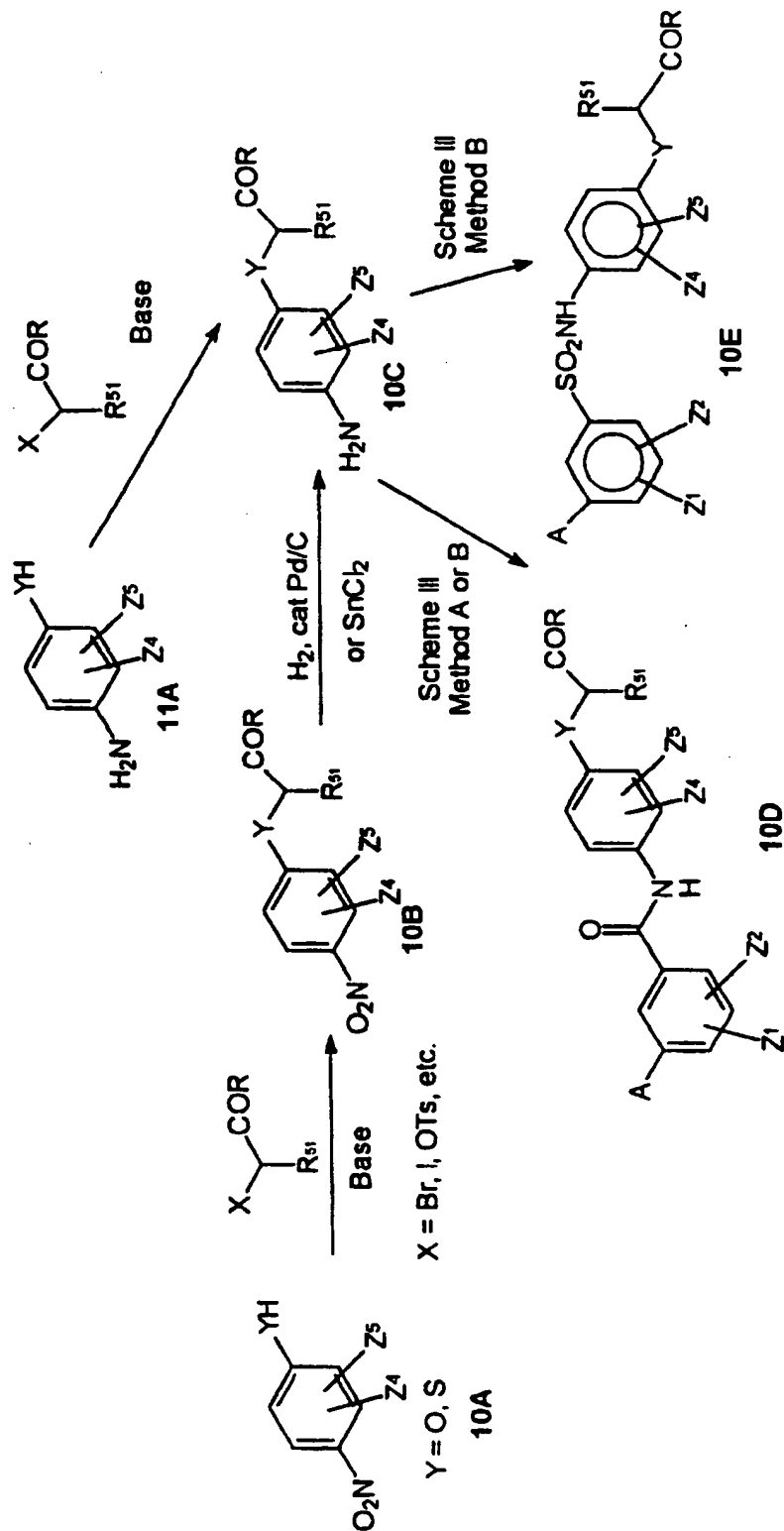
- 51 -

Scheme IV outlines methodologies for preparing tyrosine derivatives of the present invention. Thus, Scheme IV, Step 1 outlines the formation of the nitrobenzyl ether tyrosine derivative by reaction of nitrobenzyl halide or sulfonate ester with a properly protected tyrosine derivative (e.g. BOC protected) using a base such as potassium carbonate and solvent such as acetone. The BOC protecting group can be removed using standard procedures (TFA, or HCl/EtOAc, or HCl/dioxane) to give the primary amine, 6A. The amine can be functionalized with various electrophiles (e.g. Scheme II(A)) giving the secondary amine, 6B. The nitro group may be reduced by, among other reagents, tin (II) chloride dihydrate in EtOAc (60°C) giving after work-up the aniline, 6C. The aniline may be functionalized to the guanidino group by reaction with amidino pyrazine, to give 6D-1, or otherwise functionalized in similar fashion to Scheme I, to give 6D-2 to 6D-4. Hydrolysis either under standard basic conditions (e.g. LiOH or NaOH/H₂O/dioxane or THF) or by using an appropriate enzyme (e.g. porcine liver esterase in pH 8 buffer) gives the carboxylic acid derivative.

SCHEME V**Method A**

SCHEME V (Cont'd)**Method B**

SCHEME VI



- 55 -

Scheme V outlines methodologies for preparing further phenylpropionic acid derivatives. Palladium catalyzed Heck-type coupling of aryl halides 7A or 8A provided cinnamic acid derivatives 7B and 8B.

- 5 Reduction (H_2 , cat. Pd/C) provided phenylpropionic acid derivatives 7C, which were elaborated as shown in Scheme III to furnish 7D-1. Using the analogous sulfonyl chloride in place of the acid chloride provided 7E-1. Alternately, benzaldehydes 9A could be
- 10 condensed in an aldol-type reaction to give cinnamic acid derivatives 9B, which could be reduced and further elaborated as in Scheme III to give phenylpropionic acid derivatives 7D-2 and 7E-2.

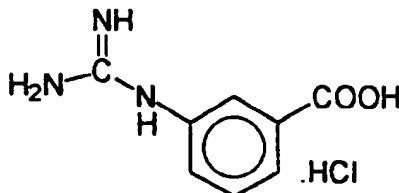
- 15 Scheme VI outlines methodologies for preparing phenoxylacetic acid and thiophenoxylacetic acid derivatives. 4-Nitrophenols or thiophenols 10A were deprotonated (NaH) and alkylated with α -haloacetic acid derivatives to give phenoxylacetic acid derivatives 10B. Reduction of the nitro group ($SnCl_2$ or H_2 , Pd/C)
- 20 provided aniline 10C. Alternately, 4-aminophenol or thiophenol 11A could be deprotonated (NaH) and alkylated with α -haloacetic acid derivatives to give phenoxylacetic acid derivatives 10C. Elaboration as shown in Scheme III gave phenoxylacetic acid derivatives
- 25 10D and 10E.

- 56 -

Example A

(3-Guanidinobenzoic acid hydrochloride)

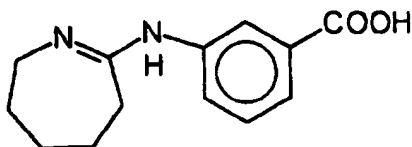
5



To 3,5-dimethylpyrazole-1-carboxamidine nitrate (6 g, 0.03 mole) (Aldrich) and diisopropylamine (3.8 g, 0.03 mole) in dioxane (20 ml) and H₂O (10 ml) was added 3-aminobenzoic acid (2.7 g, 0.02 mole). The reaction was stirred at reflux for 2.5 hours then overnight at room temperature. The resulting precipitate was filtered, washed with dioxane/H₂O and dried. The precipitate was then slurried in H₂O and acidified with concentrated HCl until a solution formed. The solvent was removed under vacuum and the residue was slurried twice in ether (ether decanted off). The product was dried under vacuum to yield 3-guanidinobenzoic acid hydrochloride (1.77 g) as a white solid. MS and NMR were consistent with the desired structure.

Example B

3-(1-Aza-2-amino-1-cycloheptenyl)benzoic acid hydrochloride



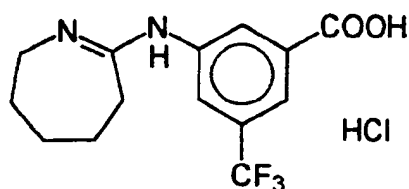
To 1-aza-2-methoxy-1-cycloheptene (3.67 g, 0.0288 mole) (Aldrich) in absolute ethanol (20 ml) was added 3-aminobenzoic acid hydrochloride (5 g, 0.0288 mole). A solution quickly formed. The reaction mixture was stirred overnight at room temperature. The resulting precipitate was filtered, washed with ether and dried under vacuum to yield 3-(1-aza-2-amino-1-cycloheptene)-benzoic acid (4.9 g).

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Example C

3-(1-aza-2-amino-1-cycloheptenyl)-5-trifluoromethylbenzoic acid hydrochloride

5



10

The title compound was synthesized according to the methodology of Example B, substituting an equivalent amount of 3-amino-5-trifluoromethyl benzoic acid [which was synthesized by reduction of 3-nitro-5-trifluoromethyl benzoic acid (Lancaster) in ethanol with 10% Pd/C under 50 psi H₂ for 4 hours] for 3-aminobenzoic acid.

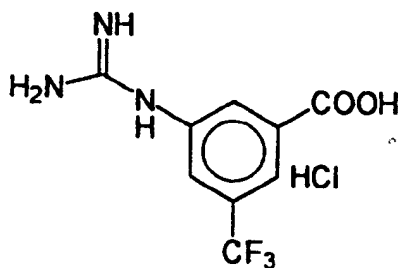
15

Example D

20

3-guanidino-5-trifluoromethylbenzoic acid, hydrochloride

25



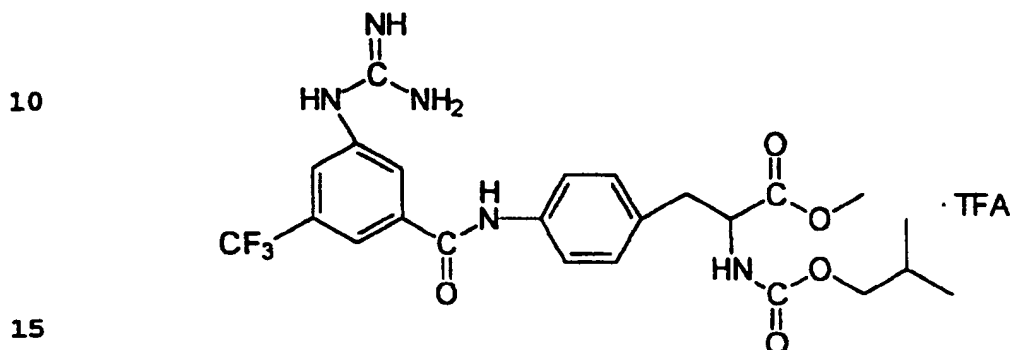
30

The title compound was synthesized according to the methodology of Example A, substituting an equivalent amount of 3-amino-5-trifluoromethylbenzoic acid (see Example C) for 3-aminobenzoic acid.

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Example 1

Synthesis of 4-[[[3-[(aminoiminomethyl)amino]-
5-(trifluoromethyl)phenyl]carbonyl]amino]-
5 N-[(2-methylpropoxy)carbonyl]phenylalanine,
methylester, trifluoroacetate salt

Step A

To p-nitrophenylalanine methyl ester hydrochloride (5 g, 0.019 mole) (Lancaster) in THF (80 mL) and triethylamine (3.88 g, 0.038 mole) was added isobutylchloroformate (2.62 g, 0.019 mole) (Sigma) dropwise at ice bath temperature. The reaction was stirred at room temperature for 3.5 hours. The resulting precipitate was filtered and washed with THF. The solvent from the filtrate was removed under vacuum. The residue was taken up in ethyl acetate. The ethyl acetate portion was worked up with saturated NaHCO₃ (1X), H₂O (1X), 1N HCl (2X), H₂O (2X), dried over MgSO₄ and removed under vacuum to yield 5.08 g of p-nitrophenylalanine methyl ester, N-isobutylcarbamate as a yellow solid.

Step B

To the product from Step A (5.08 g) in MeOH (75 mL) was added 10% Pd/C (900 mg) in a Parr bottle. This was then shaken on a Parr shaker under 50 psi H₂ at room temperature for 4.5 hours. The catalyst was

- 59 -

filtered through celite, the solvent removed under vacuum and the crude product purified by reverse phase preparatory HPLC to yield 4.5 g of p-amino-phenylalanine methyl ester, N-isobutyl carbamate,
5 trifluoroacetate as a hygroscopic yellow solid.

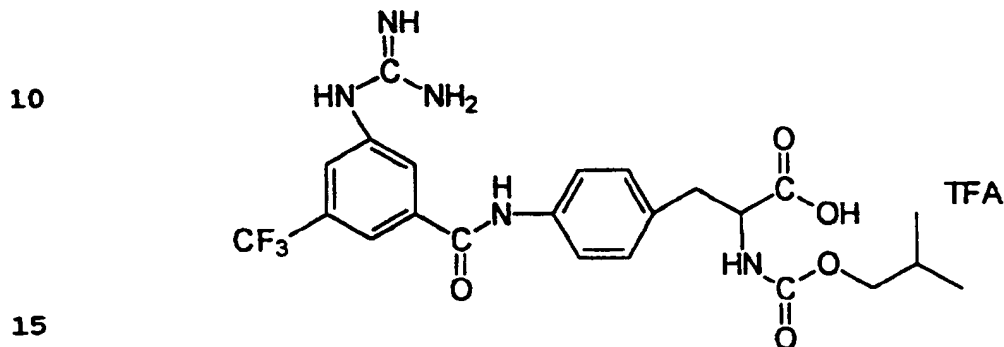
Step C

To the compound of Example D (0.43 g, 0.0015 mole) in anhydrous DMF (8 mL) and NMM (0.15 g, 0.0015 mole)
10 was added isobutylchloroformate (0.2 g, 0.0015 mole) at ice bath temperature. After stirring 5 minutes, the product from Step B (0.61 g, 0.0015 mole) in anhydrous DMF (8 mL) and NMM (0.15 g, 0.0015 mole) was added to the reaction mixture at ice bath temperature. The
15 reaction was then stirred overnight at room temperature. The solvent was removed under vacuum on a 78°C water bath and the product was isolated by reverse phase preparatory HPLC to yield (after lyophilization) 230 mg of the title compound as a white solid.
20 MS and NMR were consistent with the desired structure.

- 60 -

Example 2

Synthesis of 4-[[[3-[(aminoiminomethyl)amino]-
5-(trifluoromethyl)phenyl]carbonyl]amino]-
5 N-[(2-methylpropoxy)carbonyl]phenylalanine,
trifluoroacetate salt



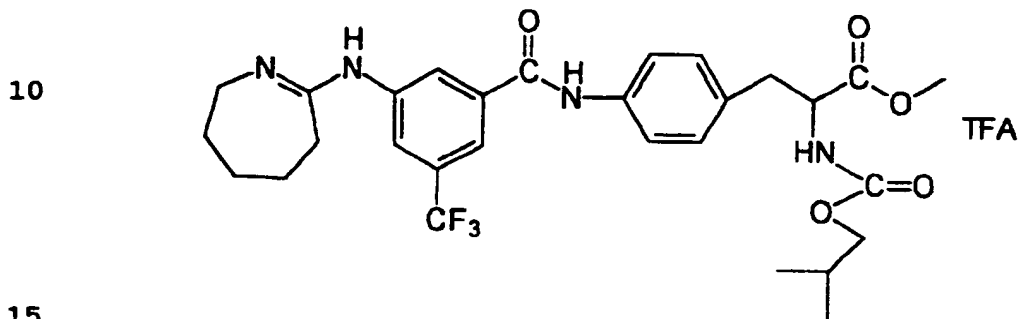
To the compound of Example 1 (200 mg, 0.0003 mole) in H₂O (10 ml) and CH₃CN (10 ml) was added LiOH (53 mg, 0.0013 mole). The reaction was stirred at room
20 temperature for 1.5 hours. The pH was lowered to 2.5 with TFA and the product was isolated by reverse phase preparatory HPLC to yield (after lyophilization) 180 mg of the title compound as a white solid.

MS and NMR were consistent with the desired
25 structure.

- 61 -

Example 3

Synthesis of N-[(2-methylpropoxy)carbonyl]-4-[[[3-
[(3,4,5,6-tetrahydro-2H-azepin-7-yl)amino]-5-
(trifluoromethyl)phenyl]carbonyl]amino]-
phenylalanine, methyl ester, trifluoroacetate salt



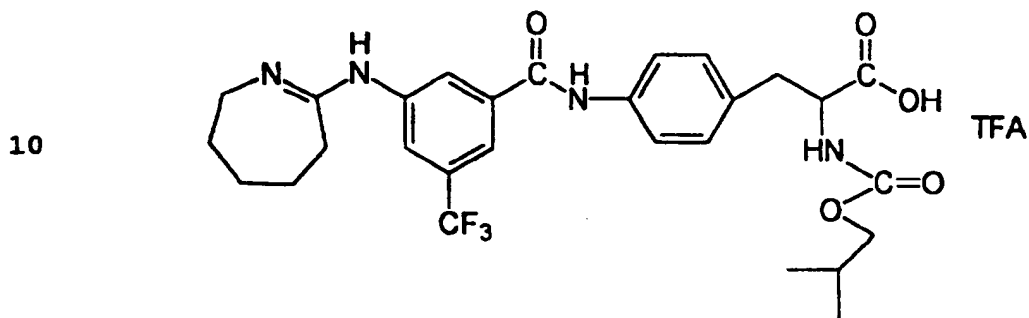
The above compound was prepared according to the methodology of Example 1, substituting an equivalent amount of the compound of Example C for the compound of Example D in Step C.

NMR and MS were consistent with the desired structure.

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Example 4

Synthesis of N-[(2-methylpropoxy)carbonyl]-
4-[[[3-[(3,4,5,6-tetrahydro-2H-azepin-7-yl)
5 amino]-5-(trifluoromethyl)phenyl]carbonyl]
amino]phenylalanine, trifluoroacetate salt



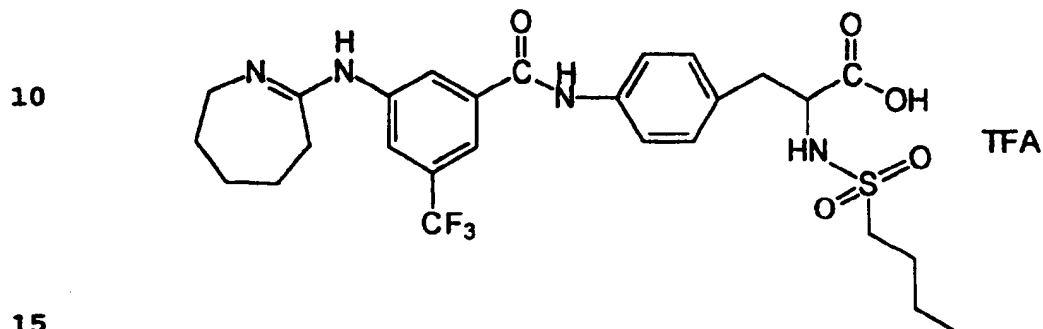
To the product from Example 3 (400 mg, 0.00058
mole) in H₂O (10 ml) and CH₃CN (10 ml) was added LiOH
(97 mg, 0.0023 mole). The reaction was stirred at room
temperature for 1 hour. The pH was lowered to 2.5 with
20 TFA and the product was isolated by reverse phase
preparatory HPLC to yield (after lyophilization) 350 mg
of the title compound as a white solid.

MS and NMR were consistent with the desired
structure.

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Example 6

Synthesis of N-(butylsulfonyl)-
4-[[[3-[(3,4,5,6-tetrahydro-2H-azepin-7-yl)-
amino]-5-(trifluoromethyl)phenyl]carbonyl]-
amino]phenylalanine, trifluoroacetate salt



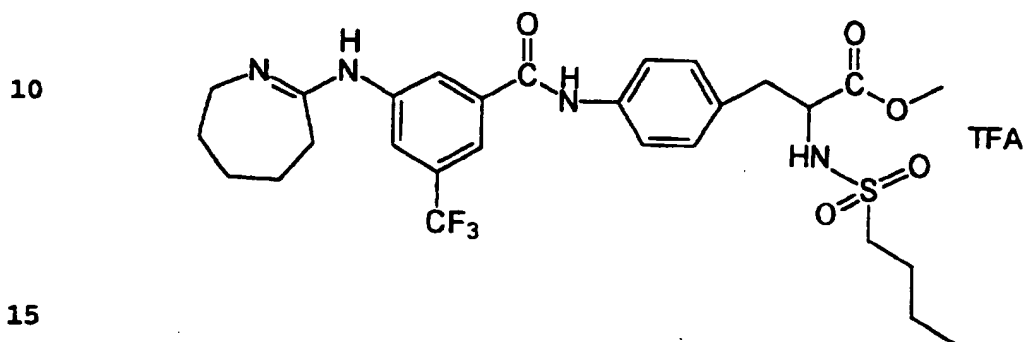
To the product of Example 5 (630 mg, 0.00089 mole) in H₂O (10 ml) CH₃CN (10 ml) was added LiOH (149 mg, 0.0035 mole). The reaction was stirred at room
20 temperature for 3.5 hours. The pH was lowered to 3.5 with TFA and the product was isolated by reverse phase preparatory HPLC to yield (after lyophilization) 560 mg of the title compound as a white solid.

MS and NMR were consistent with the desired
25 structure.

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Example 5

Synthesis of N-(butylsulfonyl)-4-[[[3-
[(3,4,5,6-tetrahydro-2H-azepin-7-yl)amino]-5-
(trifluoromethyl)phenyl]carbonyl]amino]-
phenylalanine, methyl ester, trifluoroacetate salt



The above compound was prepared according to the methodology of Example 1, substituting an equivalent amount of butanesulfonylchloride for isobutyl chloroformate in Step A and an equivalent amount of the product of Example C for the product of Example D in Step C.

NMR and MS were consistent with the desired structure.

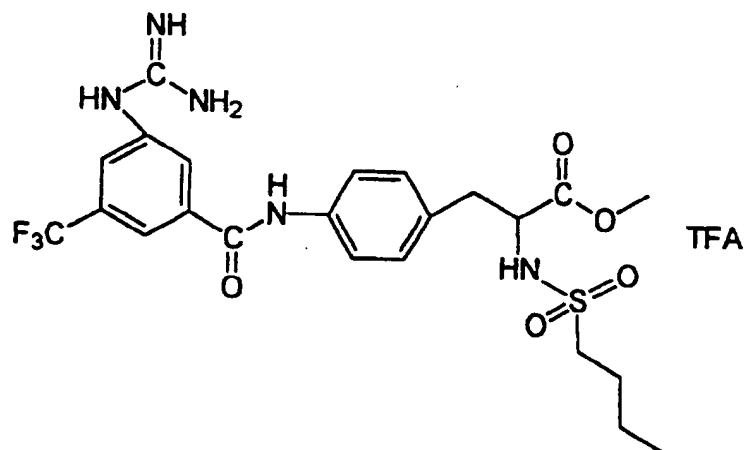
- 65 -

Example 7

Synthesis of 4-[[[3-[(aminoiminomethyl)amino]-
5-(trifluoromethyl)phenyl]carbonyl]-
amino]-N-(butylsulfonyl)phenylalanine,
methyl ester, trifluoroacetate salt

10

15



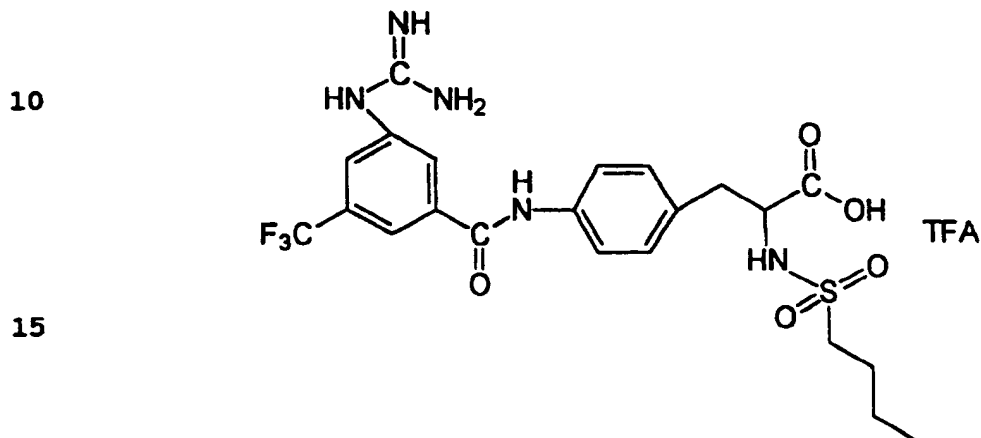
The above compound was prepared according to the methodology of Example 1, substituting an equivalent amount of butanesulfonyl chloride for isobutylchloroformate in Step A.

NMR and MS were consistent with the desired structure.

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Example 8

Synthesis of 4-[[[3-[(aminoiminomethyl)amino]-
5-(trifluoromethyl)phenyl]carbonyl]-
amino]-N-(butylsulfonyl)phenylalanine,
trifluoroacetate salt



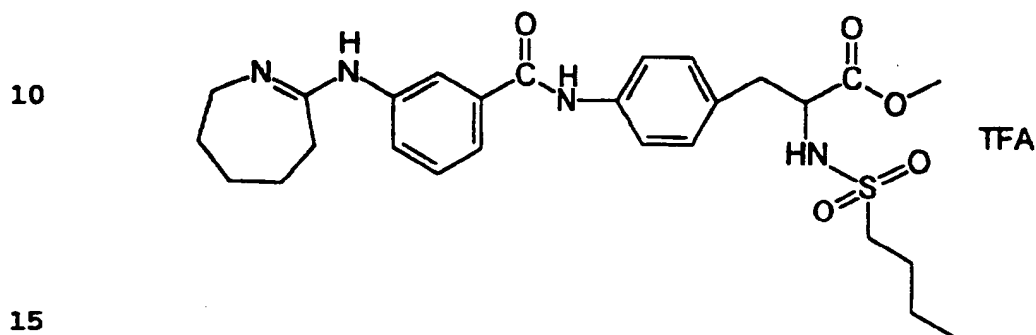
To the product of Example 7 (680 mg, 0.001 mole)
in H₂O (10 ml) and CH₃CN (10 ml) was added LiOH (174 mg,
0.004 mole). The reaction was stirred at room
temperature for 3 hours. The pH was lowered to 2.5
with TFA and the product was isolated by reverse phase
preparatory HPLC to yield (after lyophilization) 590 mg
of the title compound as a white solid.

MS and NMR were consistent with the desired
structure.

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Example 9

5 Synthesis of N-(butylsulfonyl)-4-[[[3-
 [(3,4,5,6-tetrahydro-2H-azepin-7-yl)amino]-
 phenyl]carbonyl]amino]phenylalanine,
 methyl ester, trifluoroacetate salt



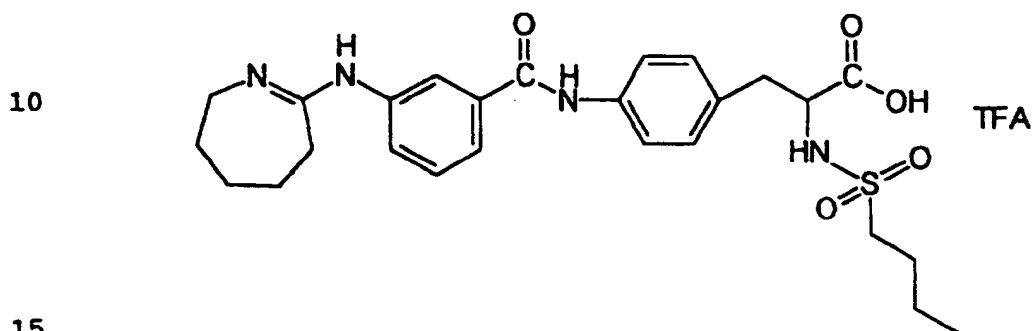
 The above compound was prepared according to the
methodology of Example 1, substituting an equivalent
amount of butanesulfonyl chloride for
20 isobutylchloroformate in Step A and an equivalent
amount of the product of Example B for the product of
Example D in Step C.

 NMR and MS were consistent with the desired
structure.

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Example 10

Synthesis of N-(butylsulfonyl)-
4-[[[3-[(3,4,5,6-tetrahydro-2H-azepin-7-yl)-
amino]phenyl]carbonyl]amino]phenylalanine,
trifluoroacetate salt



To the product of Example 9 (730 mg, 0.0011 mole) in H₂O (10 ml) and CH₃CN (10 ml) was added LiOH (191 mg, 0.0045 mole). The reaction was stirred at room
20 temperature for 2.5 hours. The pH was lowered to 2.5 with TFA and the product was isolated by reverse phase preparatory HPLC to yield (after lyophilization) 430 mg of the title compound as a white solid.

MS and NMR were consistent with the proposed
25 structure.

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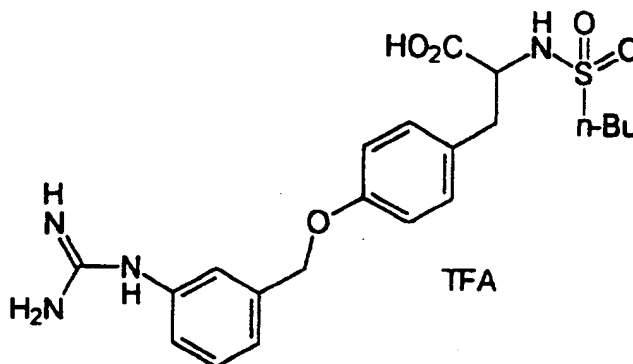
Example 11

Synthesis of O-[[3-[(aminoiminomethyl)amino]-
phenyl]methyl]-N-(butylsulfonyl)tyrosine,
trifluoroacetate salt monohydrate

5

10

15

Step A

N-t-Boc-tyrosine ethyl ester (6.0 g, 0.0194 mole) was dissolved in DMF (about 70 mL). To this solution was added, portionwise, potassium hydride (0.86 g, 0.021 mole, hexane washed mineral oil suspension). The reaction mixture was stirred until a solution formed. m-Nitrobenzyl bromide (4.54 g, 0.021 mole) was added and the reaction allowed to proceed over the weekend. Volatiles were removed to give a viscous oil that was taken up in EtOAc/dilute aqueous HCl. The aqueous layer was extracted a second time with EtOAc and the combined organic layers washed with saturated aqueous NaHCO₃, dried (Na₂SO₄), and volatiles removed to give a golden oil whose NMR and MS were consistent with the desired product.

Step B

The unpurified product from Step A was dissolved in methylene chloride (about 50 mL) and trifluoroacetic acid (20 mL) was added. Stirring was commenced and continued until no further gas evolution was noted (about 0.5 hour). Volatiles were removed until a dark

- 70 -

oil was obtained. The oil was treated with EtOAc and dilute aqueous HCl and separated. The organic layer was washed several times with dilute aqueous HCl and the acid layers were combined. The aqueous layer was
5 made basic by addition of solid NaHCO_3 and extracted (3X) with EtOAc. The pH was adjusted to about 10 by addition of 2.5 N NaOH and the aqueous layer washed with EtOAc (2X). The combined organic layers from all base washes were combined, dried (Na_2SO_4), HCl gas
10 bubbled through (until the solution was saturated) and stripped to give a tan solid (2.16 g) whose NMR and MS were consistent with the desired amino ester hydrochloride salt.

15 Step C

To the product obtained in Step B (2.16 g, 0.0062 mole), dissolved in THF (100 mL) was added triethylamine (1.38 g, 1.92 mL, 0.00136 mole) and the reaction mixture cooled to 0-5°C (ice-water/salt bath).
20 N-butanesulfonyl chloride (1.07 g, 0.0068 mole) was added in dropwise fashion. After three hours the volatiles were removed to obtain a dark oil that was worked up in similar fashion to Step A to obtain a product (2.14 g) whose NMR and MS were consistent with
25 the desired sulfonamide ester.

Step D

The product from Step C (2.14 g, 0.0046 mole) was dissolved in absolute EtOH (70 mL) and reduced to the
30 aniline using the procedure of F.D. Bellamy and K. Ou - [Tett. Let., 25, 839-842 (1984)]. Tin (II) chloride dihydrate (5.2 g, 0.023 mole) was added and the mixture heated for one hour at 60°C. The volatiles were removed to obtain a brown foam and EtOAc (100 mL) was
35 added along with saturated aqueous NaHCO_3 (200 mL) to produce a fine precipitate. The solution was clarified by passing through a celite pad. The organic layer was

- 71 -

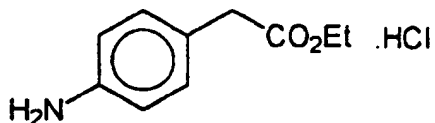
separated and the aqueous layer extracted with EtOAc (2X).

The organic layers were combined, dried (Na_2SO_4) and stripped to give a dark oil (1.75 g) that consisted
5 mostly of desired aniline as confirmed by NMR and mass spectra.

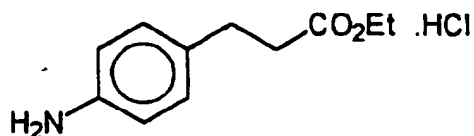
Step E

The crude aniline obtained in Step D was converted
10 to the guanidino - acid in the following manner. Aniline (1.7 g, 0.0042 mole) from Step D was dissolved in dioxane (40 mL). To this was added triethylamine (0.47 mL, 0.0047 mole), pyrazole carboxamidine (Aldrich, 0.68 g, 0.0047 mole) and water (5 mL) and the
15 reaction mixture heated to reflux. Partial conversion to product was noted after refluxing overnight. More carboxamidine was added (0.4 g) and refluxing continued. After several hours the reaction mixture was cooled and the pH adjusted to 11 by addition of
20 dilute aqueous LiOH. The reaction was maintained at pH 11 for several hours until most of the ester had hydrolyzed. The reaction mixture was made acidic by addition of TFA and the desired guanidino acid isolated by preparatory rphpic. The appropriate fractions were
25 combined and lyophilized to obtain a hydroscopic powder (0.25 g) that was determined to be the desired compound by NMR and MS.

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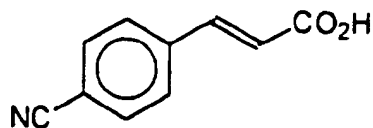
Example E

4-Aminophenylacetic acid (3 g, 19.8 mmol) was dissolved in dry ethanol (60 mL) at 0°C and a stream of hydrogen chloride gas was bubbled into the solution for 15 minutes. The solvent was removed under reduced pressure to give desired product.

Example F

20 The above compound was prepared under conditions similar to Example E, replacing 4-aminophenylacetic acid with p-aminohydrocinnamic acid.

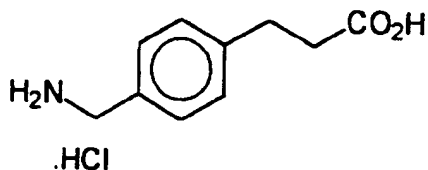
- 73 -

Example BF

5

A mixture of p-cyanobenzaldehyde (5.025 g, 38 mmol), malonic acid (4.407 g, 42 mmol), and pyridine (0.50 mL, 6.5 mmol) in absolute ethanol (10 mL) was heated to 100°C (bath) under argon. Upon heating, the mixture became a solution; and after 20-30 minutes, a yellow precipitate crashed out of solution. The reaction was monitored by TLC (10% MeOH/CH₂Cl₂). After 21.5 hours, the reaction mixture was allowed to cool to room temperature and the yellow precipitate collected by vacuum filtration. The solid was slurried with hot EtOH and collected by filtration to give the desired product as a yellow solid (4.77 g, 73% yield). NMR was consistent with proposed structure.

20

Example BG

25

The compound of Example BF (2.038 g, 11.7 mmol) was dissolved in a MeOH (15 mL)/NH₄OH (7.5 mL) mixture and hydrogenated with W-2 Raney Ni in a Parr Shaker (60 psi, 25°C) for 2.5 hours. The catalyst was filtered off and the purple filtrate concentrated in vacuo. The green solid residue was dissolved in 1 M HCl and concentrated in vacuo to give a white/green solid. The solid was purified by slurrying with 9:1 CH₃CN/MeOH mixture. The white solid was collected by vacuum

30

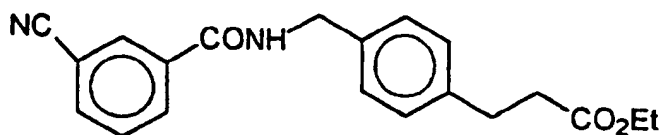
35

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Example 12

Synthesis of ethyl 4-[[[3-(cyano)phenyl]-
carbonyl]amino]methyl]benzenepropanoate

5



10 A solution of 3-cyanobenzoic acid (0.447 g, 3.0 mmol) and 1-methyl piperidine (0.37 mL, 3.0 mmol) in CH_2Cl_2 (15 mL) was cooled to 0°C . Isobutylchloroformate (0.39 mL, 3.0 mmol) was added slowly under argon and the reaction stirred for another 5 minutes. A solution
15 of the compound of Example BH (0.705 g, 2.9 mmol) and 1-methyl piperidine (0.37 mL, 3.0 mmol) in CH_2Cl_2 (3 mL) was then added and the ice bath immediately removed. The reaction was allowed to stir at room temperature for 2 hours. The reaction was concentrated *in vacuo* to
20 give a green solid residue. The residue was partitioned between EtOAc (25 mL) and water (25 mL). The organic layer was collected, washed with 1M HCl (1X25 mL), saturated NaHCO_3 (1X25 mL), and brine (1X25 mL), and then dried over MgSO_4 . Concentration *in vacuo*
25 gave the crude product as a pale yellow oil (1.17 g). The product was purified by column chromatography (50 g silica gel, 2% MeOH/ CH_2Cl_2) to give a yellow/white solid (0.476 g, 44% yield). NMR was consistent with the proposed structure.

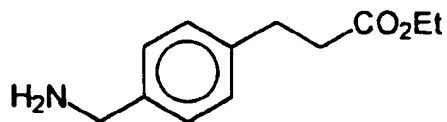
30

- 74 -

filtration to give the desired product (0.824 g, 33% yield). NMR was consistent with proposed structure.

Example BH

5



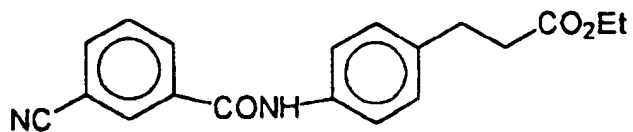
.HCl

10

A mixture of the compound in Example BG (0.824 g) in absolute EtOH (50 mL) was cooled to 0°C and HCl (g) was bubbled into it for 20 minutes. The resulting green/blue solution was allowed to stir for 2 hours. An aliquot was removed and concentrated in vacuo. ¹H NMR showed the reaction to be complete. The reaction was concentrated in vacuo to give a slightly green-tinted white solid (0.841 g). NMR was consistent with proposed structure.

20

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Example G

5

The above compound was synthesized under conditions similar to Example 12, replacing the compound of Example BH with the compound of Example F.

10

Analysis Calculated for C₁₉H₁₈N₂O₃:

C, 70.79; H, 5.63; N, 8.69.

Found: C, 70.49; H, 5.65; N, 8.62.

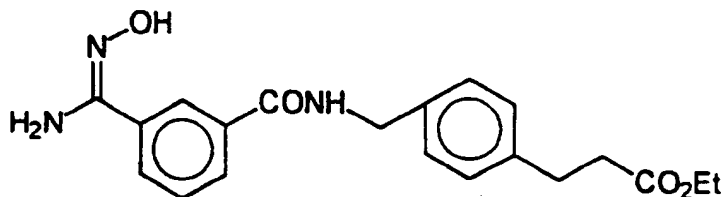
- 77 -

Example 13

Ethyl 4-[[[3-[amino(hydroxyimino)methyl]-
phenyl]carbonyl)amino]methyl]benzenepropanoate

5

10



A solution of the compound of Example 12 (0.476 g, 1.3 mmol), hydroxylamine hydrochloride (0.089 g, 1.3 mmol), and triethylamine (0.18 mL, 1.3 mmol) in absolute EtOH (10 mL) was heated to reflux (86-90°C). After 5 hours, TLC [1:1 EtOAc/hexane (10 mL) + 5 drops of AcOH] showed that starting material was still present. Additional hydroxylamine hydrochloride (0.04 g, 0.6 equivalent) and triethylamine (0.09 mL) was added. After 40 minutes, the TLC showed no change. The reaction was concentrated *in vacuo* and the residue was dissolved in H₂O (30 mL). The aqueous layer was extracted with EtOAc (40 mL). The organic layer was collected, dried over MgSO₄, and concentrated *in vacuo* to give a white solid (0.49 g). NMR was consistent with the proposed structure.

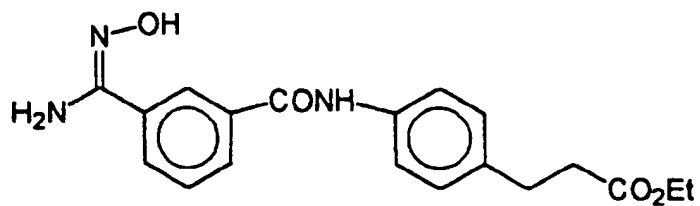
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Example 14

Ethyl 4-[[[3-[amino(hydroxyimino)methyl]phenyl]-
carbonyl]amino]benzenepropanoate

5

10



The above compound was synthesized under
conditions similar to Example 13, replacing the
15 compound of Example 12 with the compound of Example G.

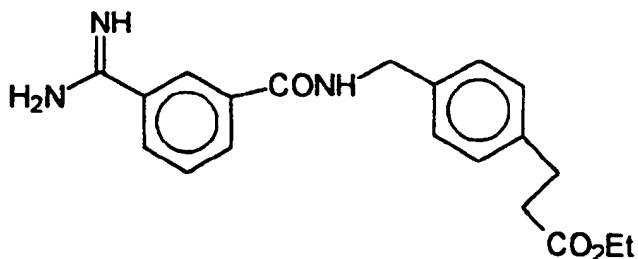
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Example 15

Ethyl 4-[[[3-(aminoiminomethyl)phenyl]-
carbonyl]amino]methyl]benzenepropanoate

5

10



The compound of Example 13 (0.49 g, 1.3 mmol) was dissolved in AcOH and hydrogenated with 4% Pd/C (53% wet, 0.050 g) in a Parr Shaker (60 psi, 60C). The catalyst was filtered off and the filtrate concentrated in vacuo to give a white solid. The solid was slurried with acetonitrile and the resulting white solid was collected by vacuum filtration (0.423 g).

20 Analysis Calculated for $C_{20}H_{23}N_3O_3 \cdot 1.6$ AcOH:

C, 61.99; H, 6.59; N, 9.35.

Found: C, 61.81; H, 6.50; N, 9.42.

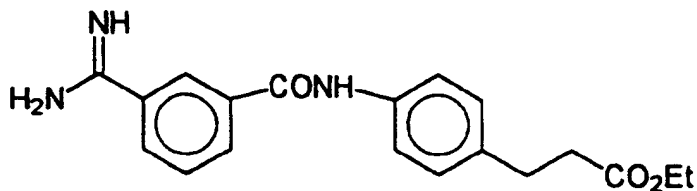
$M^+ = 353$.

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Example 17

Ethyl 4-[[[3-(aminoiminomethyl)phenyl]-
carbonyl]amino]benzenepropanoate

5



10

The compound of Example 14 was reduced under
conditions similar to the conditions described in
Example 15, replacing the compound of Example 13 with
15 the compound of Example 14.

Analysis Calculated for $C_{19}H_{31}N_3O_3 \cdot 1.7 \text{ AcOH}$:

C, 60.94; H, 6.35; N, 9.52.

Found: C, 61.02; H, 6.38; N, 9.12.

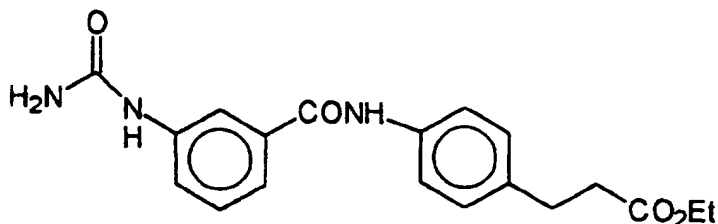
- 85 -

Example 22

Ethyl 4-[[[3-[(aminocarbonyl)amino]phenyl]-
carbonyl]amino]benzenepropanoate

5

10



A mixture of the compound of Example H (480 mg, 1.54 mmol) and acetic acid (1 mL) in water (2 mL) was heated to 38°C (bath). A solution of potassium cyanate (250 mg, 3.08 mmol) in water (2 mL) was then added slowly. The reaction became cloudy and a white precipitate resulted. The reaction was allowed to cool to room temperature and stirred for 1.5 hours. The reaction was monitored by TLC (10% MeOH/CH₂Cl₂). The white solid was collected by vacuum filtration and washed with water (0.469 g, 58% yield). The product was purified by column chromatography (10% MeOH/CH₂Cl₂) to give 150 mg desired compound.

25 Analysis Calculated for C₁₉H₂₁N₃O₄ + 0.1 H₂O:

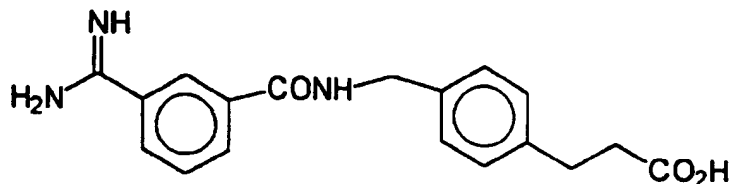
C, 63.89; H, 5.98; N, 11.76.

Found: C, 63.66; H, 5.52; N, 11.56.

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Example 19

Synthesis of 4-[[[3-(aminoiminomethyl)phenyl]carbonyl]amino]methyl]benzenepropanoic acid, trifluoroacetate salt



To a mixture of the compound of Example 15 (0.200 g, 0.57 mmol) in 1 M phosphate buffer (8 mL) was added esterase from porcine liver (Sigma, 0.5 mL) at room temperature. The reaction was stirred for 18 hours and then concentrated in vacuo. A solution of 1 M HCl (3 mL)/CH₃CN (3 mL) was added to the resulting residue and the undissolved solid filtered off. The filtrate was collected, concentrated in vacuo, and purified by HPLC-Method 1 to give the desired product as a white solid (0.17 g).

Analysis Calculated for C₁₈H₁₉N₃O₃·1.0 TFA+0.1 H₂O:

C, 54.45; H, 4.61; N, 9.52.

Found: C, 54.47; H, 4.53; N, 9.52.

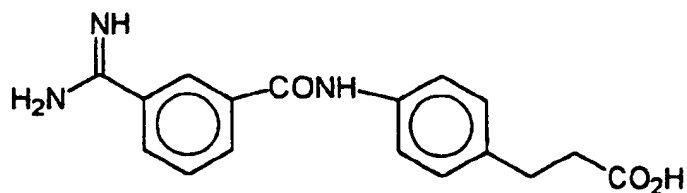
MH⁺=326.

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Example 20

4-[[[3-(Aminoiminomethyl)phenyl]carbonyl]-
amino]benzenepropanoic acid

5



10

The above compound was synthesized under conditions similar to the conditions described in Example 19 replacing the compound of Example 15 with the compound of Example 17.

15

Analysis Calculated for $C_{17}H_{17}N_3O_3 \cdot 1TFA$:

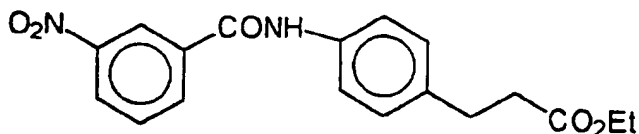
C, 53.65; H, 4.27; N, 9.88.

Found: C, 53.41; H, 4.17; N, 9.56.

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Example GA

5



10 A solution of 3-nitrobenzoic acid (0.5 g, 3 mmol) and 1-methyl piperidine (0.37 mL, 3 mmol) in CH_2Cl_2 (15 mL) was cooled to 0°C and isobutyl chloroformate (0.4 mL, 3 mmol) was added under argon. The reaction was allowed to stir for 5 minutes before adding a mixture of the compound of Example F (0.687 g, 3 mmol) and 1-methyl piperidine (0.37 mL, 3 mmol) in CH_2Cl_2 (3 mL).
15 The ice bath was removed and the reaction was allowed to stir at room temperature over 24 hours. The reaction was concentrated in vacuo and the residue was purified by column chromatography (300 g silica gel, 1% MeOH/ CH_2Cl_2) to give the desired product as a yellow
20 solid.

Analysis Calculated for $\text{C}_{18}\text{H}_{18}\text{N}_2\text{O}_5$:

C, 63.15; H, 5.30; N, 8.18.

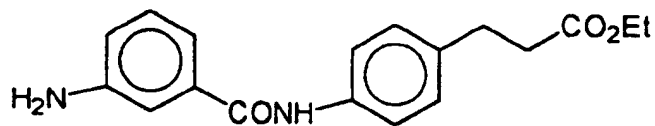
Found: C, 63.14; H, 5.41; N, 8.13.

25

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Example H

5



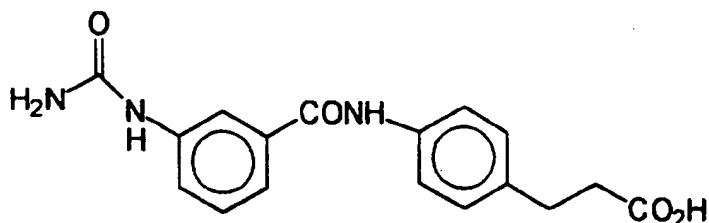
The compound of Example GA (1 g, 2.9 mmol) was hydrogenated (4% Pd/C, EtOH-THF, 5 psi, room temperature, 1.5 hours) and the filtrate concentrated in vacuo to give a yellow oil (0.8 g, 88% yield). NMR was consistent with the proposed structure.

10

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Example 24

Synthesis of 4-[[[3-[(aminocarbonyl)amino]phenyl]carbonyl]amino]benzenepropanoic acid



The Compound of Example 22 (0.12 g, 0.34 mmol) was dissolved in MeOH (2 mL) and 1 M LiOH (1 mL) was added. The reaction was stirred at room temperature over 16 hours. The reaction was concentrated *in vacuo* to give a white solid. The solid was dissolved in a small amount of H₂O and acidified with 1 drop of TFA. The mixture was concentrated *in vacuo* and the residue was purified by HPLC - Method 1 to give a white solid (0.06 g,).

Analysis Calculated for C₁₇H₁₇N₃O:

C, 62.38; H, 5.24; N, 12.84.

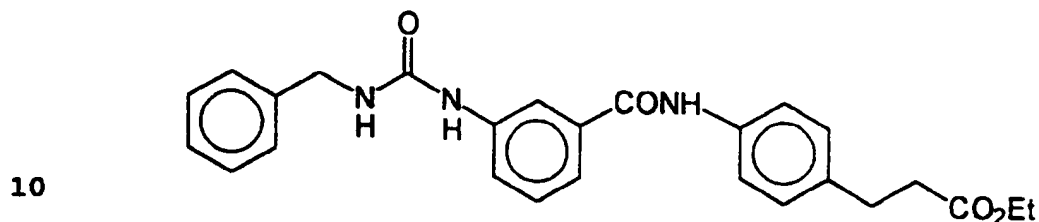
Found: C, 62.00; H, 5.53; N, 12.75.

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Example 26

Synthesis of ethyl 4-[[[3-[[[(phenylmethyl)amino]carbonyl]amino]phenyl]carbonyl]amino]benzenepropanoate

5



To a solution of benzyl isocyanate (0.146 g, 1.1 mmol) in CH_2Cl_2 (6 mL) was added a solution of the compound of Example H (0.36 g, 1.15 mmol) in CH_2Cl_2 (2 mL) under argon. The flask containing the compound of Example H was rinsed with CH_2Cl_2 (1 mL) and added to the reaction. The reaction was stirred at room temperature for 72 hours. The reaction was concentrated *in vacuo* and ether added to the yellow oil. Upon addition, the oil solidified. The resulting white solid was collected by vacuum filtration and washed with a small amount of ether (0.3 g).

Analysis Calculated for $\text{C}_{26}\text{H}_{27}\text{N}_3\text{O}_4$:

25

C, 70.10; H, 6.11; N, 9.43.

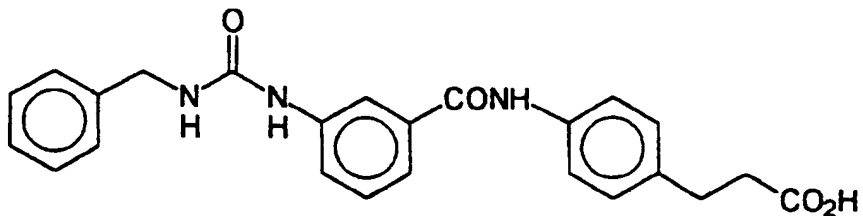
Found: C, 70.12; H, 6.35; N, 9.45.

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Example 28

Synthesis of 4-[[[3-[[[(phenylmethyl)amino]carbonyl]amino]phenyl]carbonyl]amino]benzenepropanoic acid

5



10

The compound of Example 26 was hydrolyzed under conditions similar to the conditions described in Example 24 to provide the title compound.

15 Analysis Calculated for $C_{24}H_{23}N_3O_4 \cdot 0.6 H_2O$:

C, 67.31; H, 5.70; N, 9.81.

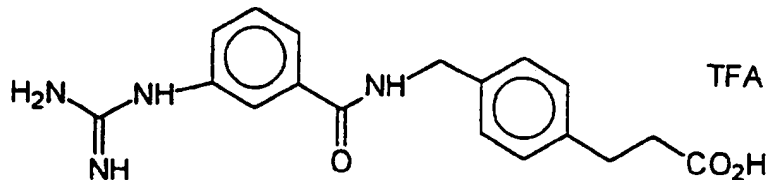
Found: C, 67.24; H, 5.35; N, 9.80.

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Example 29

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]amino]
methyl]benzenepropanoic acid, trifluoroacetate salt

5



10

A solution of the compound of Example A (0.216 g, 1.0 mmol) and 1-methyl piperidine (0.12 mL, 1.0 mmol) in DMF (5 mL) was cooled to 0°C and isobutyl chloroformate (0.13 mL, 1.0 mmol) was added under argon. The reaction was allowed to stir for 5 minutes before adding a mixture of the compound of Example BG (0.216 g, 1.0 mmol) and 1-methyl piperidine (0.12 mL, 1.0 mmol) in DMF (2 mL). The flask containing the compound of Example BG was rinsed with DMF (1 mL) and the rinse added to the reaction. The ice bath was removed after addition and the reaction was allowed to stir at room temperature over 24 hours. The reaction was concentrated in vacuo and the residue purified by HPLC-Method 1 to give the desired product as a white solid (0.051 g).

25

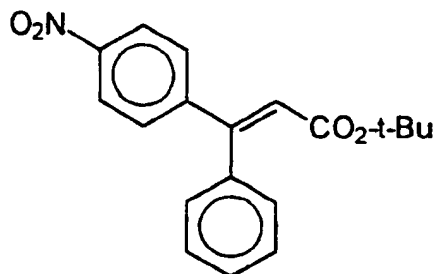
Analysis Calculated for $C_{18}H_{20}N_4O_3 \cdot 0.3 H_2O$:

C, 52.24; H, 4.73; N, 12.18.

Found: C, 52.35; H, 4.64; N, 12.21.

30 MH+=341.

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Example J

5

10 A suspension of 60% NaH in mineral oil (washed with hexane before use, 0.54 g, 13.5 mmol) in distilled THF (15 mL) was cooled to 0°C and t-butyl P,P-dimethylphosphononacetate (2.7 mL, 13.5 mmol) was added very slowly under argon. Vigorous bubbling was observed and the reaction became a white slurry. The

15 reaction was allowed to stir at 0°C for 1.5 hours before adding a solution of 4-nitrobenzophenone (3.073 g, 13.2 mmol) in THF (15 mL). The flask containing 4-nitrobenzophenone was rinsed with THF (5 mL) and the solution added to the reaction. The reaction was

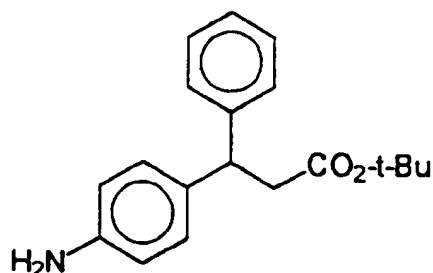
20 allowed to warm to room temperature. After 2 hours, the reaction was quenched with water (25 mL) and extracted with EtOAc (2X25 mL). The organic layers were collected, dried over MgSO₄, and concentrated in vacuo to give a pale yellow solid (4.87 g). The crude

25 material was purified by column chromatography (150 g silica gel, 5% EtOAc/hexane) and one isomer was isolated as a white solid (1.97 g). NMR was consistent with proposed structure.

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Example K

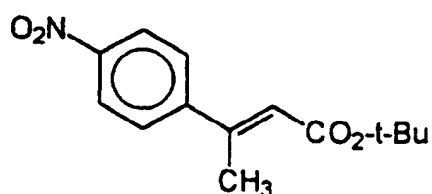
5



10 The compound of Example J (1.97 g, 6.15 mmol) was hydrogenated (EtOH/THF, 4% Pd/C, 5 psi, room temperature, 24 hours) and the filtrate concentrated in vacuo. The residue was purified by column chromatography (70 g silica, 1:1 hexane/EtOAc) to give the desired product as a white solid (1.51 g, 83%
15 yield). NMR was consistent with the proposed structure.

Example L

20



25 4-Nitroacetophenone (1.45 g, 8.8 mmol) was treated with t-butyl P,P-dimethylphosphonoacetate (2 g, 9 mmol) under similar conditions to the conditions described in Example J. The crude compound was chromatographed on silica gel using 10% EtOAc/Hex as eluant to give 700 mg of pure desired compound.

30

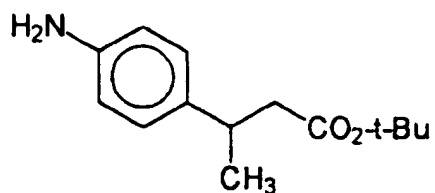
Analysis Calculated for $C_{14}H_{17}NO_4$:

C, 63.87; H, 6.51; N, 5.32.

Found: C, 63.79; H, 6.21; N, 5.16

35

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Example M

5

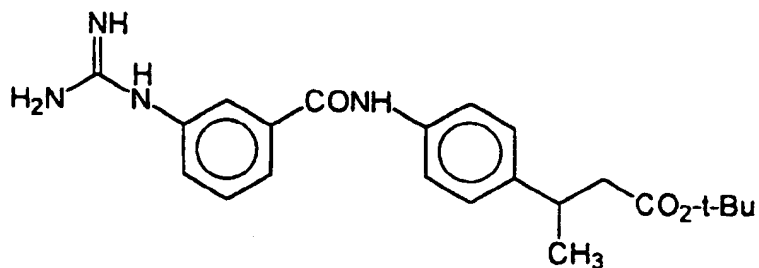
The compound of Example L was hydrogenated
(EtOH/THF, 4% Pd/C, 5 psi, room temperature, 1 hour)
10 and the filtrate concentrated *in vacuo* to afford 600 mg
of the desired compound as brown oil.

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Example 30

Synthesis of 1,1-dimethylethyl 4-[[[3-
[(aminoiminomethyl)amino]phenyl]carbonyl]amino]-
5 β -methylbenzenepropanoate, trifluoroacetate salt

10



15

The compound of Example M was coupled with the
compound of Example A under similar reaction conditions
as described in Example 34.

20

Analysis Calculated for $C_{22}H_{28}N_4O_3 \cdot 1.5 \text{ TFA} \cdot 1.1 \text{ H}_2\text{O}$:

C, 51.12; H, 5.44; N, 9.54.

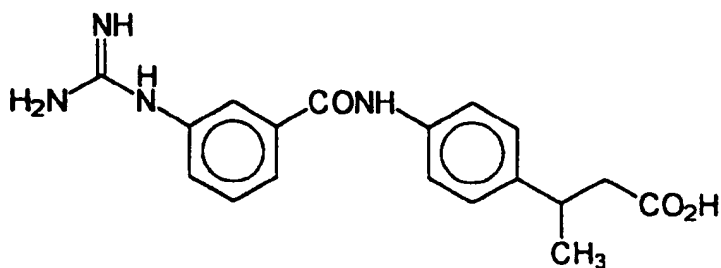
Found: C, 51.12; H, 5.05; N, 9.59.

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Example 31

Synthesis of 4-[[[3-[(aminoiminomethyl)amino]phenyl]
carbonyl]amino]- β -methylbenzenepropanoic acid,
5 trifluoroacetate salt

10



15 The compound of Example 30 was hydrolyzed with TFA
under conditions similar to those described in Example
33.

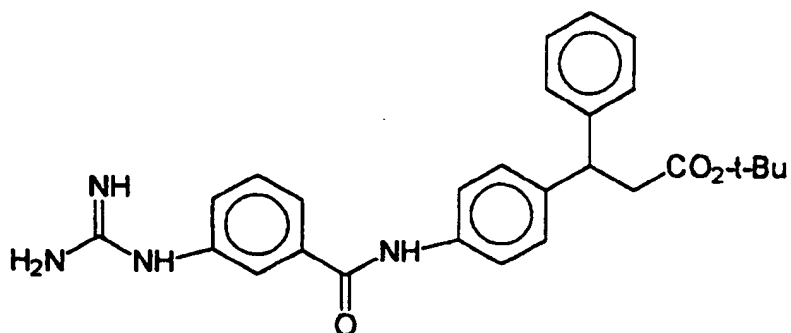
20 Analysis Calculated for $C_{18}H_{20}N_4O_3 \cdot 1.2 \text{ TFA} + 0.8 \text{ H}_2\text{O}$:
C, 49.84; H, 4.67; N, 11.40.
Found: C, 49.99; H, 4.33; N, 11.44.

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Example 32

1,1-Dimethylethyl 4-[[[3-[(aminoiminomethyl)-
amino]phenyl]carbonyl]amino] β -
phenyl benzenepropanoate

10



15

The compound of Example K (0.511 g, 1.7 mmol) was coupled with the compound of Example A under the same reaction conditions as described in Example 34. The crude material was purified by HPLC-Method 1 to give the desired product as a pale yellow oil (0.270 g, 35%
20 yield).

Analysis Calculated for $C_{27}H_{30}N_4O_3 \cdot 1.1 \text{ TFA} + 1.0 \text{ H}_2\text{O}$:

C, 58.26; H, 5.54; N, 9.31.

Found: C, 58.00; H, 5.19; N, 9.67.

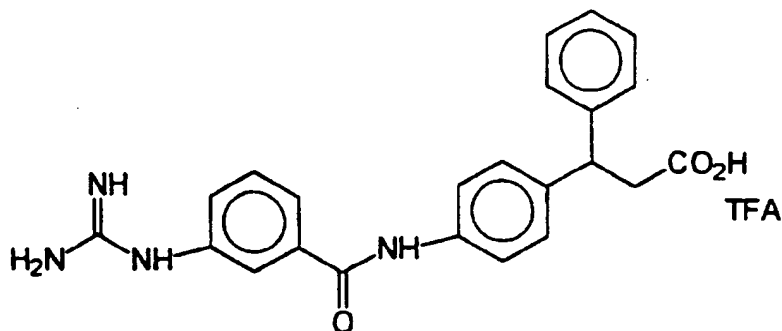
25 $M^+ = 458$.

- 96 -

Example 33

Synthesis of 4-[[[3-[(aminoiminomethyl)amino]phenyl]
carbonyl]amino]- β -phenylbenzenepropanoic acid,
5 trifluoroacetate salt

10



15 A solution of the compound of Example 32 (0.23 g)
in CH_2Cl_2 (4 mL) was cooled to 0°C and TFA (3 mL) was
added under argon. The ice bath was removed and the
reaction allowed to warm to room temperature. The
reaction was stirred for 4.5 hours, then concentrated
20 under a stream of H_2 . The residue was purified by
HPLC-Method 1 to give the desired product as a white
solid (0.085 g).

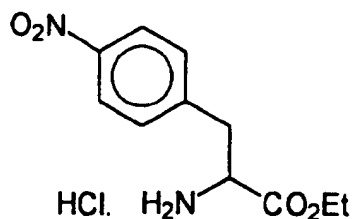
Analysis Calculated for $\text{C}_{23}\text{H}_{22}\text{N}_4\text{O}_3 \cdot 1.0 \text{ TFA} + 0.5 \text{ H}_2\text{O}$:

25 C, 57.14; H, 4.60; N, 10.66.

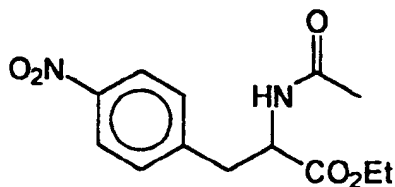
Found: C, 57.27; H, 4.65; N, 10.69.

$\text{MH}^+ = 403$.

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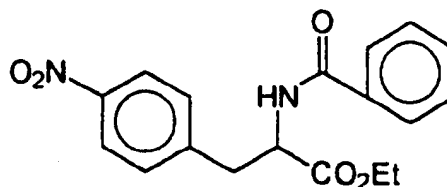
Example N

10 4-Nitro-DL-phenylalanine (5 g, 23.8 mmol) was
suspended in dry ethanol (80 mL) at 0°C and a stream of
hydrogen chloride gas was bubbled into the solution for 10
minutes. The mixture was then refluxed overnight. The
reaction was cooled to room temperature and the solvent
removed under reduced pressure to give 6.3 g of the
15 desired compound as white solid.

Example O

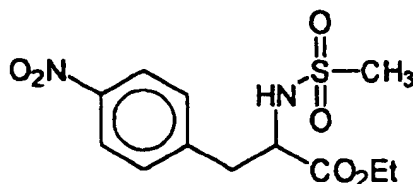
25 4-Nitro-DL-phenylalanine hydrochloride ethyl ester as
prepared in Example N (1.5 g, 5.5 mmol) was dissolved in
methylene chloride (30 mL) and cooled to 0°C. To this was
added acetyl chloride (475 mg, 6 mmol) and trimethylamine
(1.7 mL, 12 mmol). The reaction mixture was warmed to room
temperature and stirred overnight. The solvent was
30 removed under reduced pressure. Water was added and the
solution was extracted with ethyl acetate. The combined
organic layers were washed with brine, dried over Na₂SO₄,
filtered and concentrated in vacuo to give 1.5 g of the
desired compound as white solid.

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Example P

5

The above compound was prepared in the same manner as described in Example O, replacing acetyl chloride with benzoyl chloride. The residue was chromatographed on silica gel using EtOAc/Heptane (20/80) as eluant to give 1.7 g of the desired compound as yellow solid.

Example Q

15

20

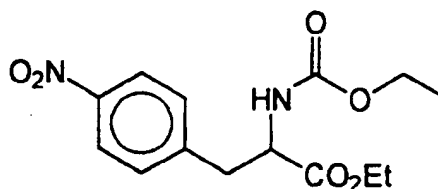
The above compound was prepared in the same manner as described in Example O, replacing acetyl chloride with methane sulfonylchloride. The residue was chromatographed on silica gel using EtOAc/Heptane (50/50) as eluant to give 1.2 g of the desired compound as a white solid.

25

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Example R

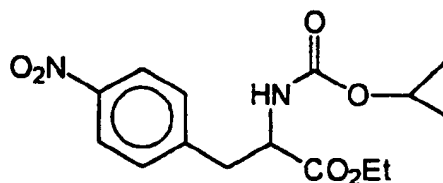
5



10 The above compound was prepared in the same manner as described in Example O replacing acetyl chloride with ethyl chloroformate. The residue was chromatographed on silica gel using EtOAc/Heptane (50/50) as eluant to give 600 mg of the desired compound as yellow oil.

Example S

15



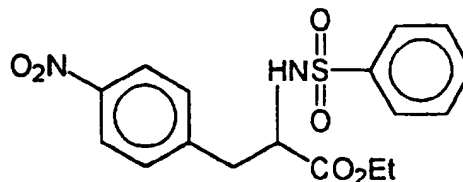
20

25 The above compound was prepared in the same manner as described in Example O, replacing acetyl chloride with isopropyl chloroformate. The residue was chromatographed on silica gel using EtOAc/Heptane (50/50) as eluant to give 600 mg of the desired compound.

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Example T

5

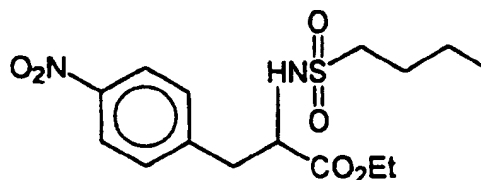


10 The above compound was prepared in the same manner as described in Example O, replacing acetyl chloride with benzenesulfonylchloride. The residue was chromatographed on silica gel using EtOAc/Hexane (50/50) as eluant to give 1.3 g of the desired compound as white solid.

15

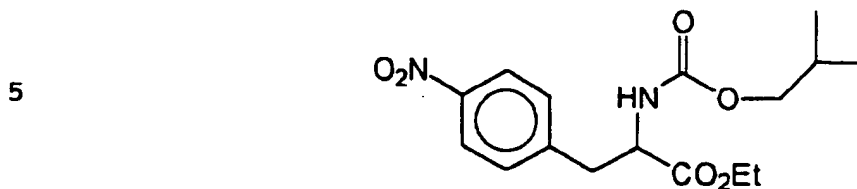
Example U

20



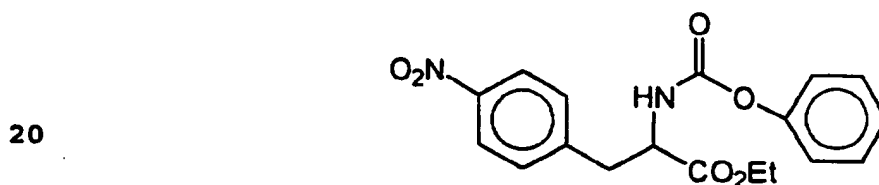
25 The above compound was prepared in the same manner as described in Example O, replacing acetyl chloride with 1-butananesulfonyl chloride. The residue was chromatographed on silica gel using EtOAc/Heptane (50/50) as eluant to give 900 mg of the desired compound.

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Example V

10 The above compound was prepared in the same manner as described in Example O, replacing acetyl chloride with isobutyl chloroformate. The residue was chromatographed on silica gel using EtOAc/Heptane (50/50) as eluant to give 1.1 g of the desired compound.

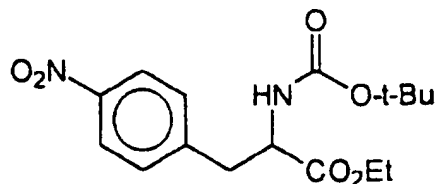
15

Example W

25 The above compound was prepared in the same manner as described in Example O, replacing acetyl chloride with phenyl chloroformate. The residue was chromatographed on silica gel using EtOAc/Heptane (50/5) as eluant to give 1.1 g of the desired compound.

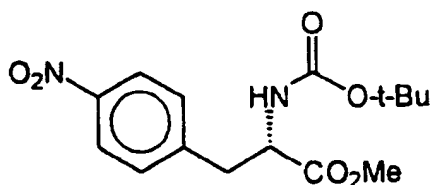
30

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Example X

5

A mixture of the compound of Example N (1.5 g 5.5 mmol), di-tert-butyl dicarbonate (1.32 g, 6 mmol) and potassium carbonate (2.3 g, 16.5 mmol) in THF/H₂O (1:1, 20 mL) was stirred at room temperature overnight. The solution was concentrated and the residue was dissolved in ethyl acetate. The solution was treated with water, and extracted with ethyl ether. The combined organic phases were washed with brine, dried over Na₂SO₄ and evaporated to afford 1.8 g yellow solid.

Example Y

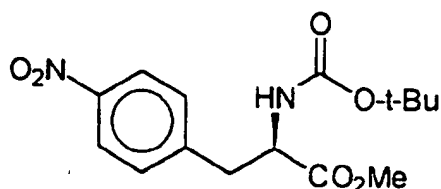
20

A mixture of (S)-4-nitrophenylalanine methyl ester hydrochloride (1.43 g, 5.5 mmol), di-tert-butyl dicarbonate (1.3 g, 6 mmol) and potassium carbonate (2.3 g, 16.5 mmol) in THF/H₂O (1:1, 20 mL) was stirred at room temperature overnight. The solution was concentrated and the residue was dissolved in ethyl acetate. The solution was treated with water, and extracted with ethyl ether. The combined organic phases were washed with brine, dried over Na₂SO₄ and evaporated to afford 1.8 g yellow solid.

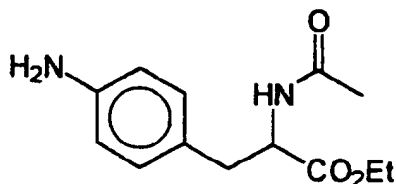
25

30

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Example Z

5
10
15
N-Boc-p-nitro-D-phenylalanine (1 g, 3.2 mmol, Bachem) was treated with ethereal CH_2N_2 solution (15 mL), (prepared from 1.65 g N-nitroso-N-methyl urea) at 0°C . The reaction mixture was warmed to room temperature and stirred for 3 hours. The solvent was removed under reduced pressure to give 1.1 g title compound as white solid.

Example AA

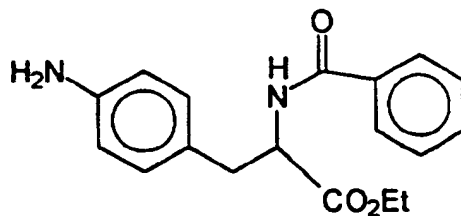
20
25
30
The product of Example O (1.5 g, 5.3 mmol) was dissolved in EtOH/THF and transferred to a Parr Shaker containing a catalytic amount of 4% Pd/C. The reaction was shaken for 16 hours at room temperature under 5 psi pressure of H_2 . The reaction mixture was filtered and concentrated to afford 1.2 g of a brown solid.

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The following compounds were prepared in the same manner as described in Example AA.

Example AB

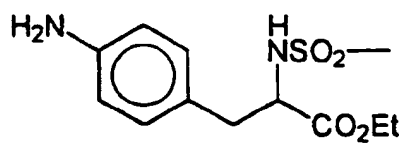
5



10

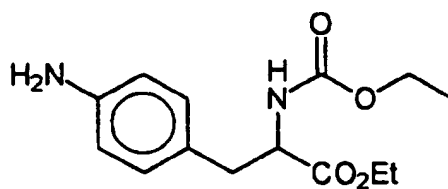
Example AC

15



Example AD

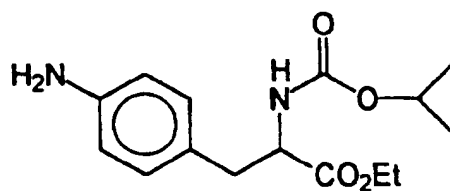
20



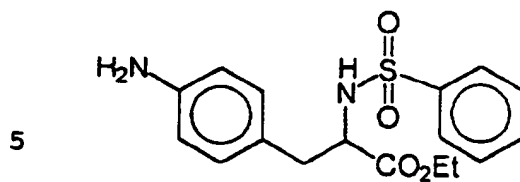
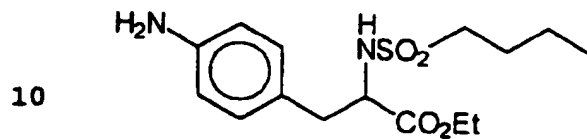
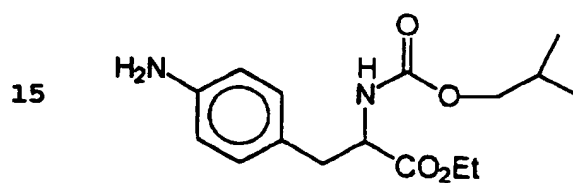
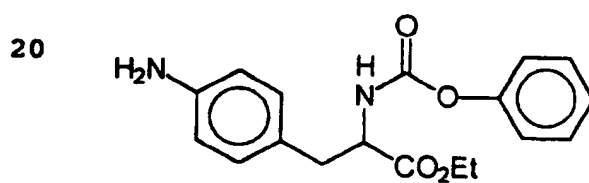
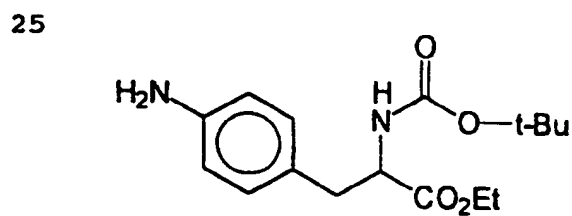
25

Example AE

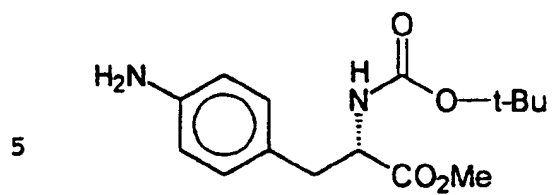
30



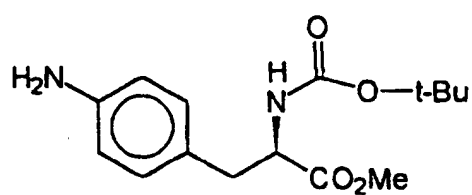
- 105 -

Example AFExample AGExample AHExample AIExample AJ

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Example AKExample AL

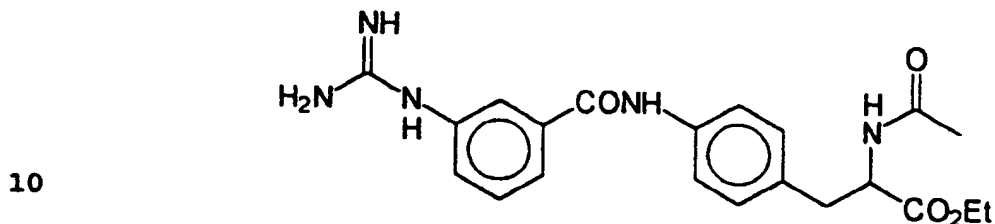
10



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Example 34

Synthesis of N-acetyl-4-[[[3-[(aminoiminomethyl)amino]
phenyl)carbonyl]amino]phenylalanine ethyl ester,
5 trifluoroacetate salt



To a stirred solution of the compound of Example A
(517 mg, 2.4 mmol) in dimethyl formamide (10 mL) at 0°C
was added 1-methylpiperidine (238 mg, 2.4 mmol) followed
15 by the addition of isobutyl chloroformate (328 mg, 2.4
mmol). After 5 minutes the compound of Example AA (600
mg, 2.4 mmol) in dimethyl formamide (1 mL) was introduced.
The reaction mixture was warmed to room temperature and
stirred overnight. The solvent was removed under reduced
20 pressure, and the residue was purified by reverse phase
HPLC-Method 1 to give 600 mg white solid.

Analysis Calculated for $C_{21}H_{25}N_5O_4 \cdot 1.6TFA \cdot 0.8 H_2O$:

C, 47.78; H, 4.67; N, 11.51.

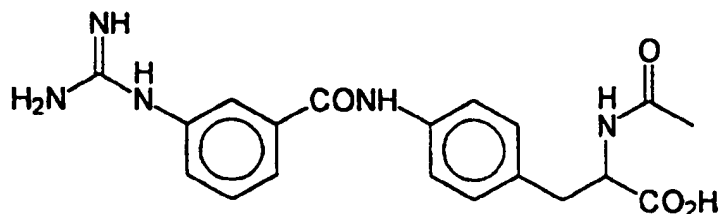
25 Found: C, 47.54; H, 4.67; N, 11.86.

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Example 35

Synthesis of N-acetyl-4-[[[3-[(aminoiminomethyl)amino]-phenyl]carbonyl]amino]phenylalanine, trifluoroacetate salt

5



10

The product of Example 34 (420 mg, 1 mmol) was dissolved in methanol (4 mL) at room temperature. Lithium hydroxide (1M, 2 mL) was added and the reaction mixture was stirred overnight. The solution was concentrated and purified by reverse phase HPLC-Method 1 to give 120 mg white solid.

15

Analysis Calculated for $C_{19}H_{21}N_5O_4 \cdot 1.6TFA \cdot 0.6 H_2O$:

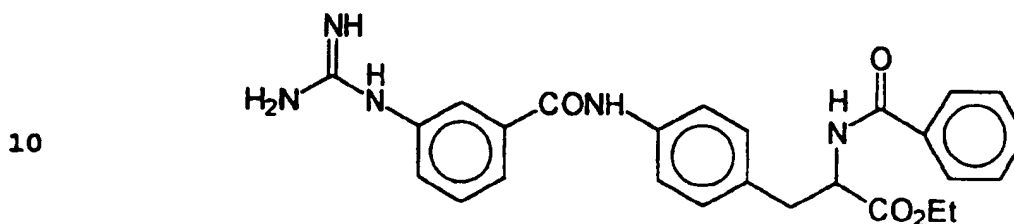
20

C, 46.24; H, 4.16; N, 12.14.

Found: C, 46.41; H, 4.15; N 11.83

Example 36

Synthesis of 4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]amino]-N-(phenylcarbonyl)phenylalanine ethyl
5 ester, monohydrate trifluoroacetate salt



The title compound was prepared in the same manner as described in Example 34, replacing the compound of Example 15 AA with the compound of Example AB.

Analysis Calculated for $C_{26}H_{27}N_5O_4 \cdot 1.1 \text{ TFA} \cdot 1 \text{ H}_2O$:

C, 54.90; H, 4.92; N, 11.35.

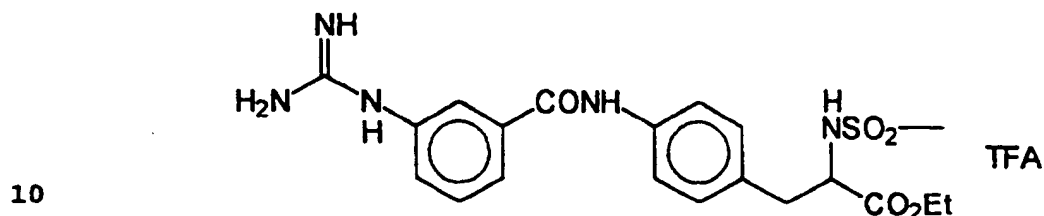
Found: C, 54.82; H, 4.60; N, 11.51.

20

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Example 37

Synthesis of 4-[[[3-[(aminoiminomethyl)amino]phenyl]
carbonyl]amino]-N-(methylsulfonyl)phenylalanine ethyl
5 ester, monohydrate trifluoroacetate salt



The title compound was prepared in the same manner as
described in Example 34, replacing the compound of Example
AA with the compound of Example AC.

15

Analysis Calculated for C₂₀H₂₅N₅O₅S·1TFA·1 H₂O:

C, 45.59; H, 4.87; N, 12.08.

Found: C, 45.38; H, 4.79; N, 11.89.

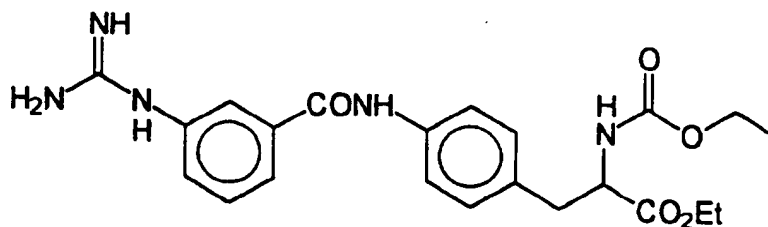
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Example 38

Synthesis of 4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]amino]-N-(ethoxycarbonyl)phenylalanine ethyl
5 ester, trifluoroacetate salt

10



15

The title compound was prepared in the same manner as
described in Example 34, replacing the compound of Example
AA with the compound of Example AD.

Analysis Calculated for $C_{22}H_{27}N_5O_5 \cdot 1.4 \text{ TFA} \cdot 0.8 \text{ H}_2\text{O}$:

C, 48.39; H, 4.91; N, 11.38.

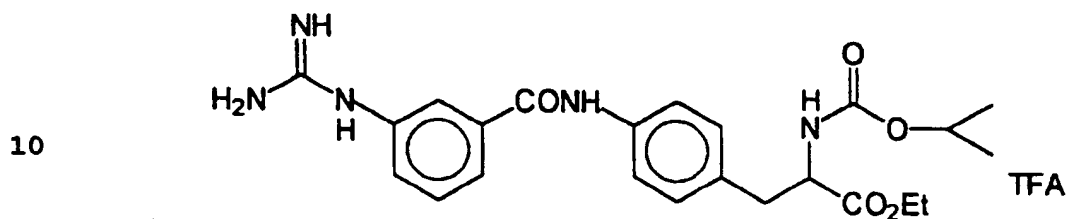
Found: C, 48.43; H, 4.98; N, 11.45.

20

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Example 39

Synthesis of 4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]amino]-N-[(1-methylethoxy)carbonyl]phenylalanine
5 ethyl ester, monohydrate trifluoroacetate salt



The title compound was prepared in the same manner as
described in Example 34, replacing the compound of Example
15 AA with the compound of Example AE.

Analysis Calculated for $C_{23}H_{29}N_5O_5 \cdot 1 \text{ TFA} \cdot 1 \text{ H}_2\text{O}$:

C, 51.11; H, 5.49; N, 11.92.

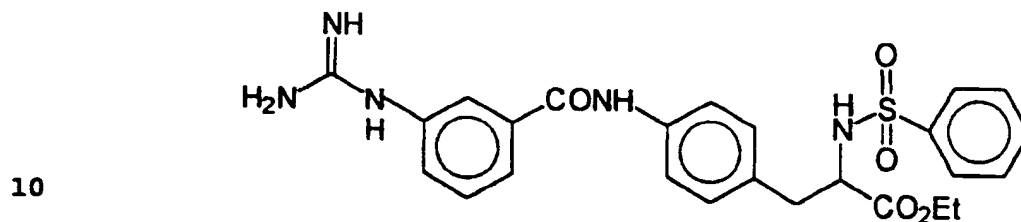
Found: C, 50.91; H, 5.26; N, 11.92.

20

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Example 40

Synthesis of 4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]amino]-N-(phenylsulfonyl)phenylalanine ethyl
5 ester, monohydrate trifluoroacetate salt



The title compound was prepared in the same manner as
described in Example 34 replacing the compound of Example
15 AA with the compound of Example AF.

Analysis Calculated for $C_{25}H_{27}N_5O_5S \cdot 1.1 \text{ TFA} \cdot 1 \text{ H}_2\text{O}$:

C, 50.03; H, 4.65; N, 10.72.

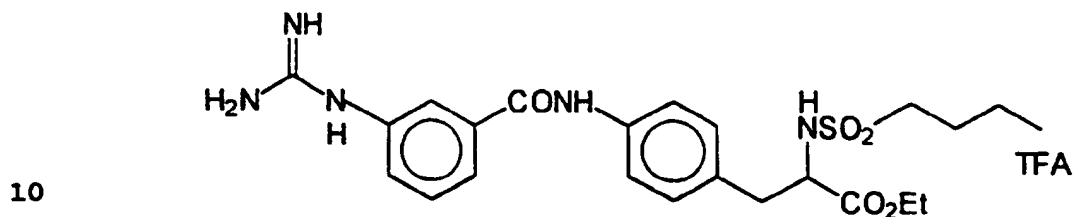
Found: C, 50.02; H, 4.33; N, 10.77.

20

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Example 41

Synthesis of 4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]amino]-N-(butylsulfonyl)phenylalanine ethyl
5 ester, trifluoroacetate salt



The title compound was prepared in the same manner as
described in Example 34, replacing the compound of Example
AA with the compound of Example AG.

15 Analysis Calculated for C₂₃H₃₁N₅O₅S·1TFA·0.2H₂O:

C, 49.45; H, 5.38; N, 11.53.

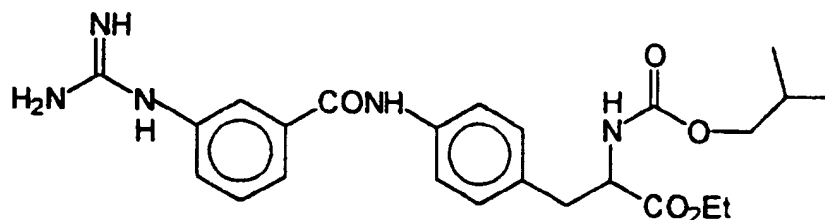
Found: C, 49.25; H, 5.04; N, 11.92.

20

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Example 42

Synthesis of 4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]amino]-N-[(2-methylpropoxy)carbonyl]phenyl-
alanine, ethyl ester, trifluoroacetate salt



The above compound was prepared in the same manner as described in Example 34, replacing the compound of Example AA with the compound of Example AH.

Analysis Calculated for $C_{24}H_{31}N_5O_5 \cdot 1.4TFA \cdot 0.3 H_2O$:

C, 50.73; H, 5.24; N, 11.30.

Found: C, 50.33; H, 5.01; N, 11.53.

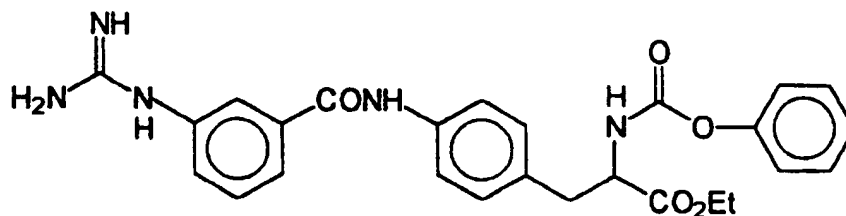
- 116 -

Example 43

4-[[[3-[(Aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(phenoxy)carbonyl]phenylalanine,
ethyl ester

5

10



The above compound was prepared in the same manner as
described in Example 34, replacing the compound of Example
AA with the compound of Example AI.

15

Analysis Calculated for $C_{26}H_{27}N_5O_4 \cdot 1.1 \text{ TFA} \cdot 1 \text{ H}_2\text{O}$:

C, 54.90; H, 4.92; N, 11.35.

Found: C, 54.82; H, 4.60; N, 11.51.

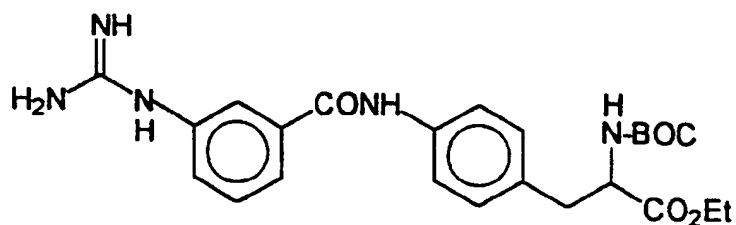
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Example 44

Synthesis of 4-[[[3-[(aminoiminomethyl)amino]
phenyl]carbonyl]amino]-N-[(1,1-dimethylethoxy)carbonyl]
5 phenylalanine ethyl ester, trifluoroacetate salt

10



15

The title compound was prepared in the same manner as
described in Example 34, replacing the compound of Example
AA with the compound of Example AJ.

Analysis Calculated for $C_{26}H_{27}N_5O_5 \cdot 1.2 \text{ TFA} \cdot 1 \text{ H}_2\text{O}$:

C, 52.94; H, 4.73; N, 10.87.

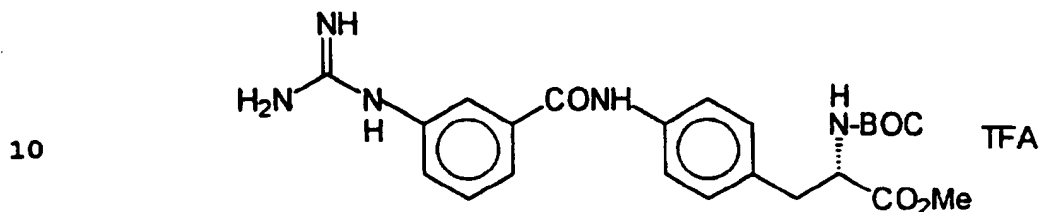
Found: C, 53.04; H, 4.71; N, 10.81.

20

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Example 45

Synthesis of 4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]amino]-N-[(1,1-dimethylethoxy)carbonyl]-L-
5 phenylalanine, methyl ester, trifluoroacetate salt



The above compound was prepared in the same manner as
described in Example 34, replacing the compound of Example
15 AA with the compound of Example AK.

Analysis Calculated for $C_{23}H_{29}N_5O_5 \cdot 1 \text{ TFA} \cdot 0.9 \text{ H}_2\text{O}$:

C, 51.26; H, 5.47; N, 11.96.

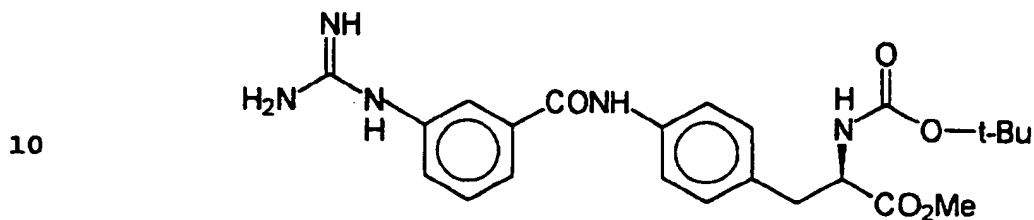
Found: C, 51.26; H, 5.26; N, 11.85.

20

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Example 46

Synthesis of 4-[[[3-[(aminoiminomethyl)amino]-
phenyl]carbonyl]amino]-N-[(1,1-dimethylethoxy)carbonyl]-D-
5 phenylalanine, methyl ester



The above compound was prepared in the same manner as
described in Example 34, replacing the compound of Example
15 AA with the compound of Example AL.

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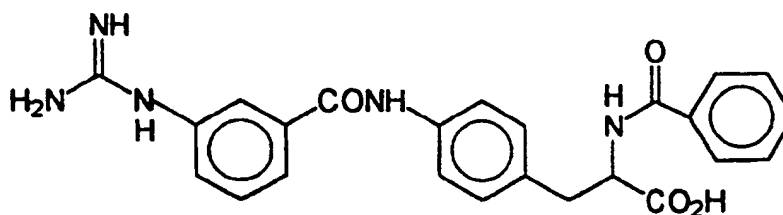
The above compounds of Examples 36 through 46 were hydrolyzed in the same manner as described in Example 35 to produce the compounds of the following Examples:

5

Example 47

Synthesis of 4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]amino]-N-(phenylcarbonyl)phenylalanine, trifluoroacetate salt

10



15

Analysis Calculated for $C_{24}H_{23}N_5O_4 \cdot 1.4 \text{ TFA} \cdot 0.9 \text{ H}_2\text{O}$:

C, 51.81; H, 4.25; N, 11.27.

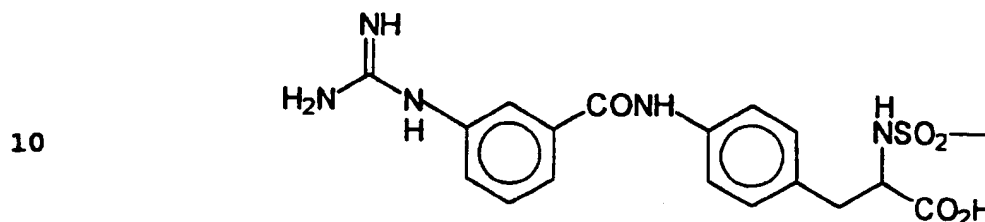
Found: C, 51.75; H, 4.25; N, 11.45.

20

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Example 48

Synthesis of 4-[[[3-[(aminoiminomethyl)amino]phenyl]
carbonyl]amino]-N-(methylsulfonyl)phenylalanine,
5 trifluoroacetate salt



Analysis Calculated for $C_{18}H_{21}N_5O_5S \cdot 1.1 \text{ TFA} \cdot 0.2 \text{ H}_2\text{O}$:

C, 44.73; H, 4.20; N, 13.04.

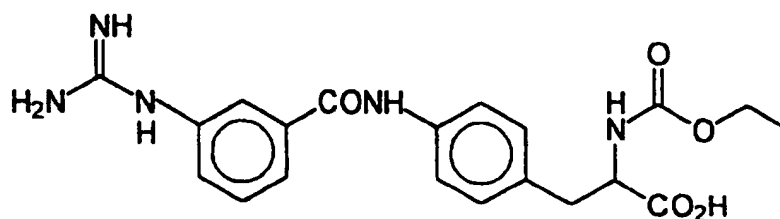
15 Found: C, 44.66; H, 4.02; N, 13.11.

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Example 49

Synthesis of 4-[[[3-[(aminoiminomethyl)amino]phenyl]
carbonyl]amino]-N-(ethoxycarbonyl)phenylalanine,
5 trifluoroacetate salt

10



Analysis Calculated for $C_{28}H_{23}N_5O_5 \cdot 1.4 \text{ TFA} \cdot 0.5 \text{ H}_2\text{O}$:

C, 46.89; H, 4.38; N, 11.99.

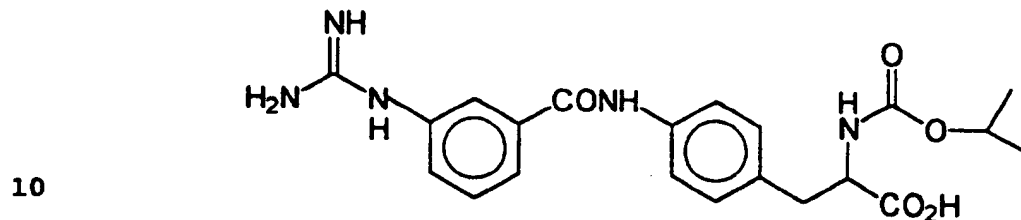
15

Found: C, 46.96; H, 4.33; N, 11.75.

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Example 50

Synthesis of 4-[[[3-[(aminoiminomethyl)amino]phenyl]
carbonyl]amino]-N-[(1-methylethoxy)carbonyl]-
5 phenylalanine, trifluoroacetate salt



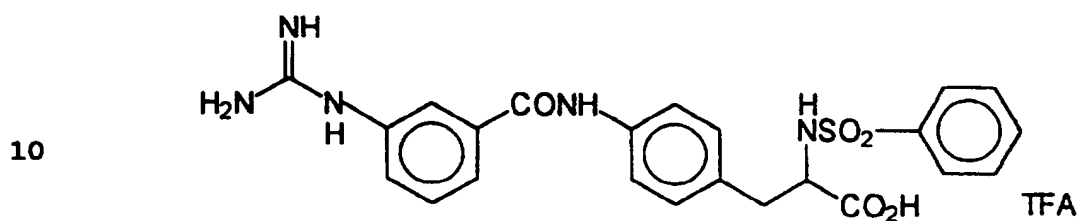
15

Analysis Calculated for $C_{21}H_{25}N_5O_5 \cdot 1.1 \text{ TFA} \cdot 0.9 \text{ H}_2\text{O}$:
C, 48.96; H, 4.97; N, 12.31.
Found: C, 48.93; H, 4.80; N, 12.48.

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Example 51

Synthesis of 4-[[[3-[(aminoiminomethyl)amino]phenyl]
carbonyl]amino]-N-(phenylsulfonyl)phenylalanine,
5 trifluoroacetate salt



Analysis Calculated for $C_{23}H_{23}N_5O_5S \cdot 1 \text{ TFA} \cdot 0.5 \text{ H}_2\text{O}$:

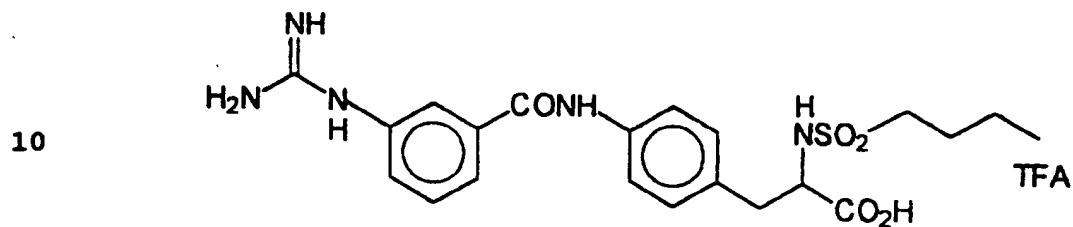
C, 49.67; H, 4.17; N, 11.58.

15 Found: C. 49.74; H, 4.22; N, 11.68.

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Example 52

Synthesis of 4-[[[3-[(aminoiminomethyl)amino]phenyl]
carbonyl]amino]-N-(butylsulfonyl)phenylalanine,
5 trifluoroacetate salt



Analysis Calculated for $C_{21}H_{27}N_5O_5S \cdot 1TFA \cdot 0.1 H_2O$:

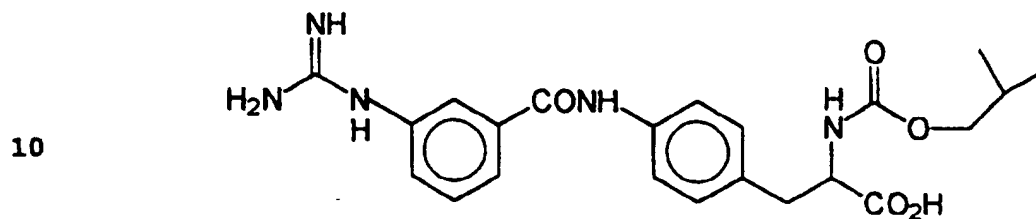
C, 47.58; H, 4.92; N, 12.13.

15 Found: C, 47.79; H, 4.84; N, 12.12.

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Example 53

Synthesis of 4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]amino]-N-[(2-methylpropoxy)carbonyl]-
5 phenylalanine, trifluoroacetate salt



Analysis Calculated for $C_{22}H_{27}N_5O_5 \cdot 1.1TFA \cdot 1H_2O$:

15 C, 54.90; H, 4.92; N, 11.35.

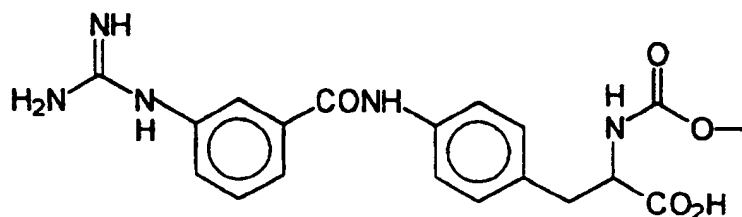
Found: C, 54.82; H, 4.60; N, 11.51.

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Example 54

Synthesis of 4-[[[3-[(aminoiminomethyl)amino]phenyl]
carbonyl]amino]-N-(methoxycarbonyl)phenylalanine,
5 trifluoroacetate salt

10



15

Analysis Calculated for $C_{19}H_{21}N_5O_5 \cdot 1.3 \text{ TFA} \cdot 0.4 \text{ H}_2\text{O}$:

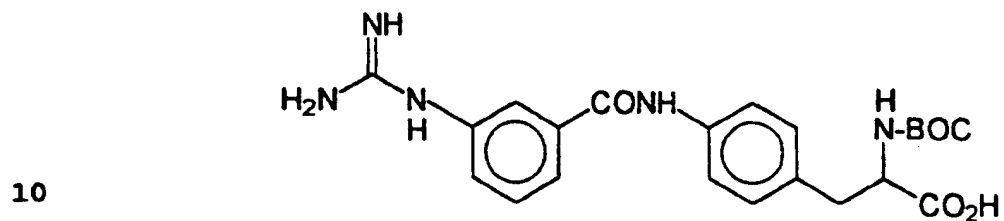
C, 46.76; H, 4.20; N, 12.62.

Found: C, 46.76; H, 3.95; N, 12.65.

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Example 55

Synthesis of 4-[[[3-[(aminoiminomethyl)amino]phenyl]
carbonyl]amino]-N-[(1,1-dimethylethoxy)carbonyl]-
5 phenylalanine, monohydrate trifluoroacetate salt



Analysis Calculated for $C_{22}H_{27}N_5O_5 \cdot 1.1TFA \cdot 1H_2O$:

C, 49.69; H, 5.19; N, 11.99.

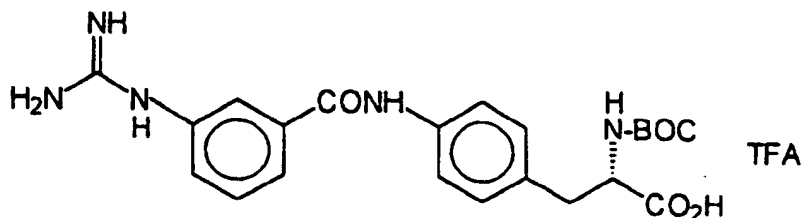
15 Found: C, 49.65; H, 4.95; N, 11.95.

- 129 -

Example 56

Synthesis of 4-[[[3-[(aminoiminomethyl)amino]phenyl]
carbonyl]amino]-N-[(1,1-dimethylethoxy)carbonyl]-
5 L-phenylalanine, trifluoroacetate salt

10



15

Analysis Calculated for $C_{22}H_{27}N_5O_5 \cdot 1 \text{ TFA} \cdot 1.1 \text{ H}_2\text{O}$:

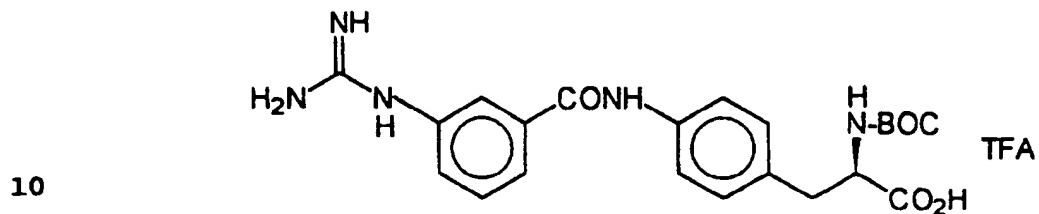
C, 50.10; H, 5.29; N, 12.17.

Found: C, 49.86; H, 5.27; N, 12.20.

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Example 57

Synthesis of 4-[[[3-[(aminoiminomethyl)amino]phenyl]
carbonyl]amino]-N-[(1,1-dimethylethoxy)carbonyl]-D-
5 phenylalanine, monohydrate trifluoroacetate salt



Analysis Calculated for $C_{22}H_{27}N_5O_5 \cdot 1 \text{ TFA} \cdot 1 \text{ H}_2\text{O}$:

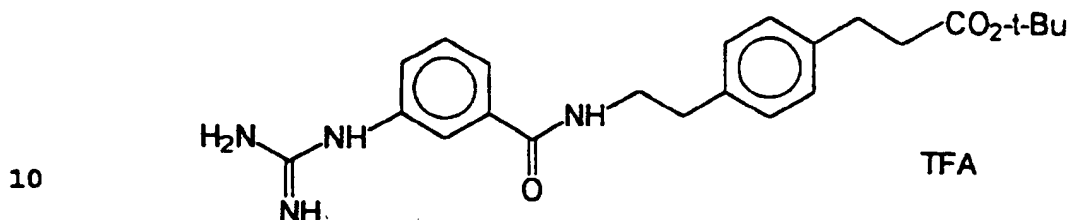
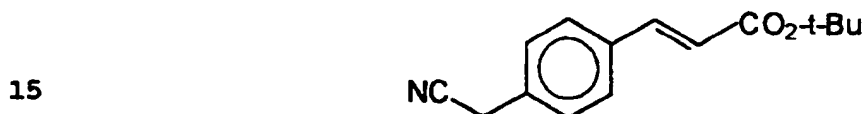
C, 50.26; H, 5.27; N, 12.21.

15 Found: C, 50.20; H, 4.95; N, 12.25.

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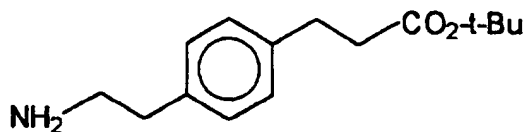
Example 58

1,1-dimethylethyl-4-[2-[[[3-[(aminoiminomethyl)amino]-
phenyl]carbonyl]amino]ethyl]benzenepropanoate,
5 monohydrate trifluoroacetate salt

Step A

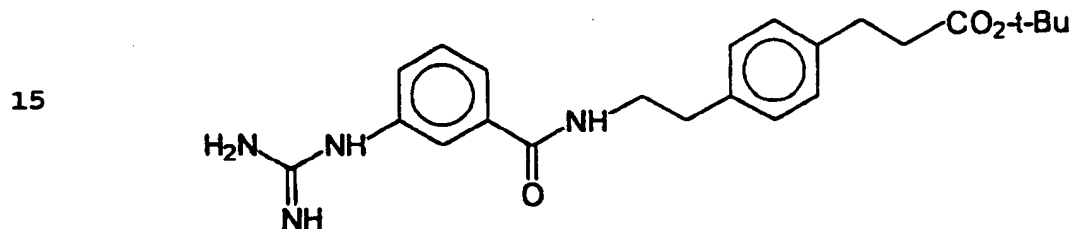
A mixture of p-bromophenylacetonitrile (5.03 g, 25.6 mmol), t-butyl acrylate (6.0 mL, 40.8 mmol), palladium acetate (0.060 g, 0.26 mmol), and tri-o-tolylphosphine (0.326 g, 1.0 mmol) in triethylamine (12 mL) was heated to
20 reflux under argon for 6 hours. The resulting orange mixture was poured into ice and acidified to pH 1 (pH paper) with 1 M HCl. The mixture was extracted with EtOAc (150 mL). The organic layer was collected, dried over
25 MgSO₄, and concentrated in vacuo to give an orange/yellow solid. The solid was recrystallized from ether/hexane to give yellow crystals (1.91 g). NMR was consistent with the proposed structure.

30 Step B



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The compound of Step A (1.9 g, 7.8 mmol) was dissolved in i-PrOH/HCl and hydrogenated with 10% Pd/C in a Parr Shaker (60 psi) for 10 hours at room temperature. The catalyst was removed and the filtrate was concentrated in vacuo. The residue was partitioned between saturated NaHCO₃ and ether. The aqueous layer was back-extracted with ether. The organic layers were combined, dried over K₂CO₃, and concentrated in vacuo to give a pale yellow oil (1.93 g, 98% yield). NMR was consistent with proposed structure.

Step C

20 The compound of Step B (0.504 g, 2.0 mmol) was coupled with the compound of Example A according to procedures described in Example 29. The crude material was purified by HPLC-Method 1 to give a sticky white solid (0.50 g).

25

Analysis Calculated for C₂₃H₃₀N₄O₁·1.0 TFA+ 1.0 H₂O:

C, 55.34; H, 6.13; N, 10.33.

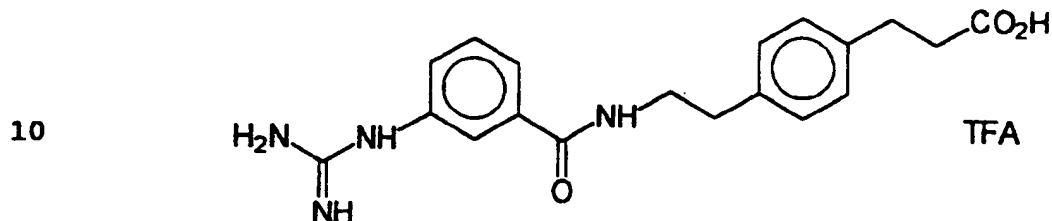
Found: C, 55.34; H, 5.77; N, 10.16. M⁺=410.

30

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Example 59

4-[2-[[[3-[(aminoiminomethyl)amino]phenyl]
carbonyl]amino]ethyl]benzenepropanoic acid,
5 trifluoroacetate salt



15 A solution of the compound of Example 58 in CH_2Cl_2 (4 mL) was cooled to 0°C and TFA (3 mL) was added under argon. The ice bath was removed and the reaction allowed to warm to room temperature. The reaction was stirred for 4.5 hours, then concentrated under a stream of N_2 . The crude material was slurried with acetonitrile and the
20 resulting white solid was collected by vacuum filtration (0.235 g).

Analysis Calculated for $\text{C}_{19}\text{H}_{22}\text{N}_4\text{O}_3 \cdot 1.0$ TFA:

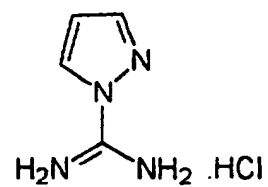
C, 53.85; H, 4.95; N, 11.96.

25 Found: C, 53.59; H, 4.93; N, 11.98. $\text{MH}^+=355$.

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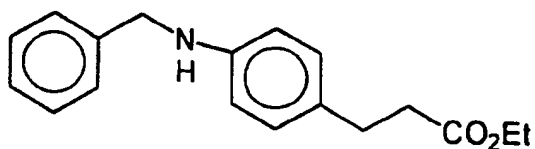
Example AM

5

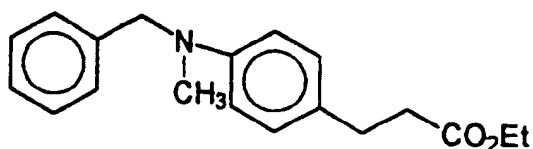


The above compound was prepared according to
10 (Bernatowicz, JOC, Vol. 57, No. 8, (1992), p. 2497-2502.
NMR was consistent with the proposed structure.

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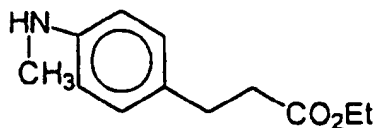
Example AN

5 To a stirred solution of benzaldehyde (763 mg, 7.2 mmol), and the compound of Example F (1.5 g, 6.5 mmol) in ethanol (10 mL) was added borane-pyridine complex (0.2 mL), and the mixture was stirred at room temperature for
10 30 minutes. The reaction mixture was then concentrated and the residue was dissolved in ethyl acetate, washed with saturated sodium bicarbonate and brine, dried over Na_2SO_4 , and evaporated. The crude product was
15 chromatographed on silica gel using EtOAc/Hexane (1:8) as eluant to give 1.3 g of the desired compound as yellow oil.

Example AO

20 To a stirred solution of formaldehyde (37%, 0.28 mL, 3.5 mmol), the compound of Example AN (0.9 g, 3.2 mmol) in ethanol (5 mL) was added borane-pyridine complex (BPC)
25 (0.32 mL 3.5 mmol), and the mixture was stirred at room temperature for 1 hour. One additional equivalent of formaldehyde and BPC were added to the reaction and stirred overnight. The reaction mixture was concentrated
30 and the residue was dissolved in ethyl acetate, washed with saturated sodium bicarbonate and brine, dried over Na_2SO_4 and evaporated. The crude product was chromatographed on silica gel using EtOAc/Hexane (1:8) as eluant to give 900 mg of the desired compound as colorless oil.

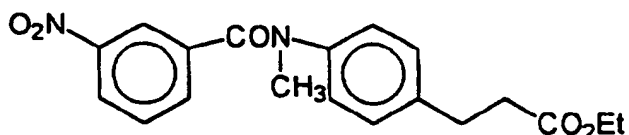
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Example AP

5

The compound of Example AO (1.3 g, 4.5 mmol) was dissolved in a mixture of EtOH (20 mL) and acetyl chloride (353 mg, 4.5 mmol) was added. This mixture was transferred to a pressure bottle with a catalytic amount of 10% Pd/C wetted with ethanol. The reaction was stirred for 3 hours at room temperature under a H₂ pressure of 5 psi. The reaction mixture was filtered and concentrated to afford 750 mg of an oil.

15

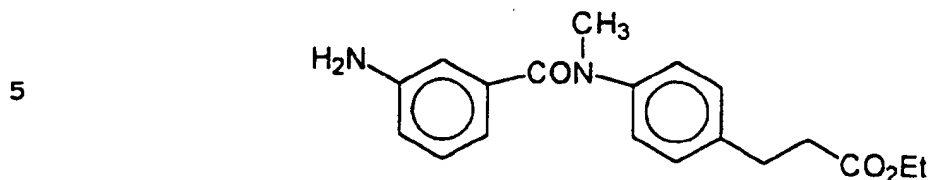
Example AO

20

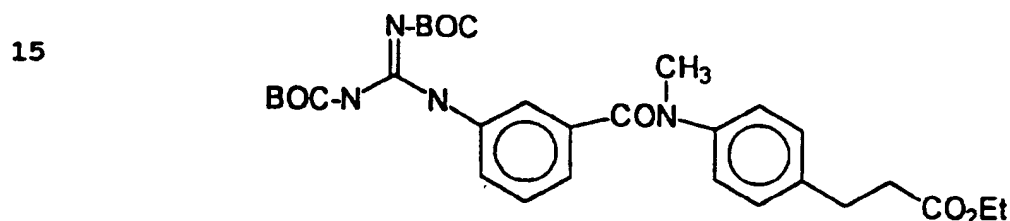
To a stirred solution of the product of Example AP (710 mg, 2.9 mmol) in methylene chloride (15 mL) at 0°C was added 3-nitrobenzoyl chloride (557 mg, 3 mmol), followed by triethylamine (0.85 mL, 6 mmol). The reaction mixture was stirred at room temperature overnight. The reaction mixture was then concentrated and the residue was dissolved in ethyl acetate, washed with water and brine, dried over Na₂SO₄ and evaporated. The residue was chromatographed on silica gel using EtOAc/Heptane (60/40) as eluant to give 1 g of the pure desired compound.

30

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Example AR

The compound of Example AQ was reduced in a similar
10 manner as described in Example AA.

Example AS

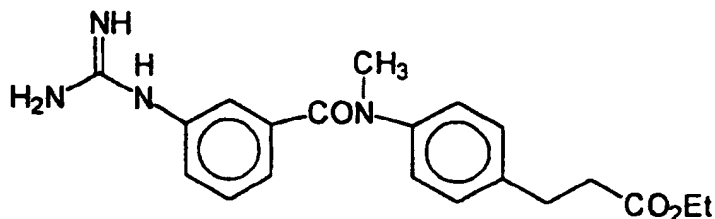
20 To a stirred mixture of the compound of Example AR
(610 mg, 1.87 mmol), the compound of Example AZ (530 mg,
1.92 mmol) and triethylamine (0.86 mL, 6.1 mmol) in
dimethyl formamide (7 mL) at 0°C under argon was added
mercury dichloride and the mixture was stirred at 0°C for
25 3 hours. The mixture was diluted with ethyl acetate and
filtered through celite. The filtrate was concentrated
and the residue was treated with water and, extracted with
ethyl acetate. The combined organic phases were washed
with brine, dried over Na₂SO₄ and evaporated. The residue
30 was chromatographed on silica gel using EtOAc/Heptane
(50/50) as eluant to give 730 mg of the pure desired
compound as a white solid.

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Example 60

Synthesis of ethyl 4-[[[3-[(aminoiminomethyl)amino]
phenyl]carbonyl]methylamino]benzenepropanoate,
5 trifluoroacetate salt

10



15

20

The compound of Example AS (720 mg, 1.27 mmol) was dissolved in methylene chloride (6 mL) and cooled to 0°C. To the solution was added trifluoroacetic acid (2 mL). After 15 minutes the ice bath was removed and the reaction stirred for 4 hours. The reaction mixture was then concentrated and the residue was purified by reverse phase HPLC-Method 1 to give 610 mg of a colorless oil.

Analysis Calculated for $C_{20}H_{24}N_4O_3 \cdot 1 \text{ TFA} \cdot 1.2 \text{ H}_2\text{O}$:

C, 52.42; H, 5.48; N, 11.11.

Found: C, 52.25; H, 5.39; N, 11.08.

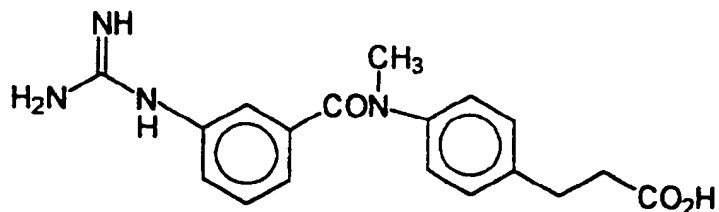
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Example 61

Synthesis of 4-[[[3-[(aminoiminomethyl)amino]phenyl]
carbonyl]methylamino]benzenepropanoic acid,

5 monohydrate trifluoroacetate salt

10



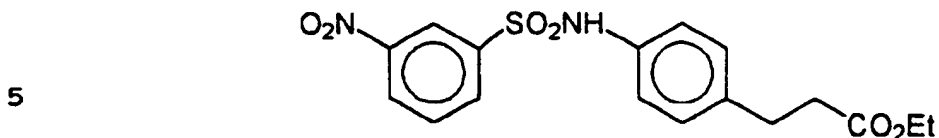
The compound of Example 60 was hydrolyzed in the same
manner as described in Example 35.

15 Analysis calculated for $C_{18}H_{20}N_4O_3 \cdot 1.6 \text{ TFA} \cdot 1 \text{ H}_2\text{O}$:

C, 47.08; H, 4.40; N, 10.36.

Found: C, 47.14; H, 4.34; N, 10.34.

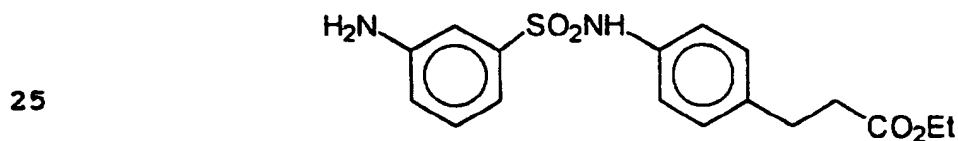
- 140 -

Example AT

To a stirred solution of the compound of Example F (1.07 g, 4.7 mmol) in methylene chloride (15 mL) at 0°C was added 3-nitrobenzenesulphonyl chloride (1.03 g, 4.7 mmol) followed by triethylamine (1.25 mL, 9 mmol). The mixture was stirred at room temperature overnight under argon. The reaction mixture was then concentrated and the residue was treated with water and extracted with

15 chloroform. The combined organic phases were washed with brine, dried over Na₂SO₄ and evaporated. The residue was chromatographed on silica gel using 1% MeOH/CH₂Cl₂ as eluant to give 1.2 g of the pure desired compound as a yellow oil.

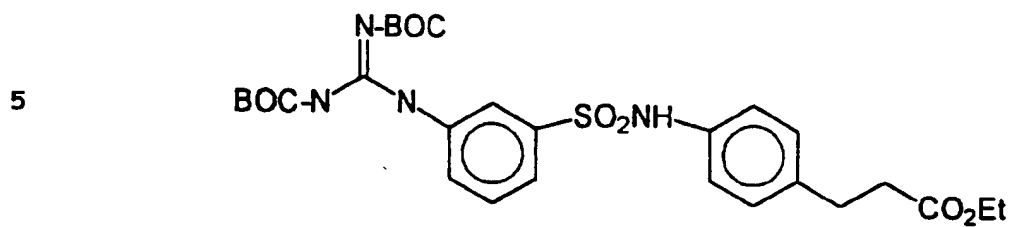
20

Example AU

The compound of Example AT was reduced in the same manner as described in Example AA.

30

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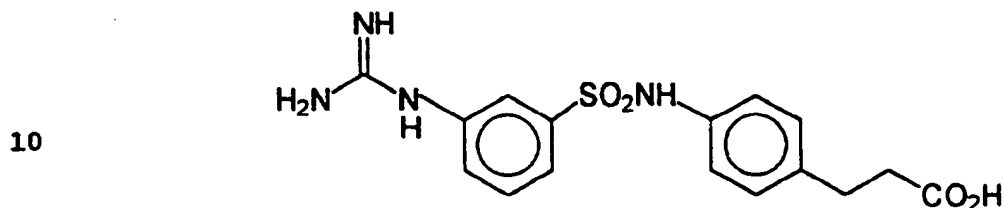
Example AV

10 The above compound was synthesized under conditions similar to those described in Example AS.

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Example 62

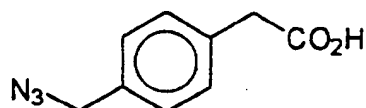
Synthesis of 4-[[[3-[(aminoiminomethyl)amino]phenyl]
sulfonyl]amino]benzenepropanoic acid,
5 trifluoroacetate salt



The compound of Example AV (130 mg, 0.22 mmol) was
dissolved in dioxane (10 mL), and treated with 6 N HCl (10
15 mL). The reaction mixture was stirred at room temperature
overnight, concentrated and purified by reverse phase
HPLC-Method 1.

Analysis Calculated for $C_{16}H_{18}N_4O_4S \cdot 1.4 \text{ TFA} \cdot 0.3 \text{ H}_2\text{O}$:
20 C, 42.81; H, 3.82; N, 10.61.
Found: C, 43.15; H, 3.62; N, 10.07.

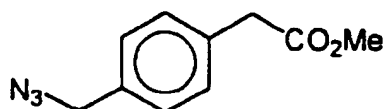
- 143 -

Example AW

5 To a stirred solution of 4-(bromomethyl)-phenylacetic acid (1.18 g, 5.15 mmol) in dimethyl formamide (10 mL) was added sodium azide (402 mg, 6.18 mmol), and mixture was heated at 60°C for 4 hours. The reaction mixture was cooled to room temperature, poured into water, and
10 extracted with ethyl acetate. The combined organic phases were washed with water and brine, dried over MgSO₄ and evaporated to afford 0.92 g of the desired compound.

Example AX

15



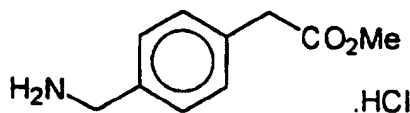
The compound of Example AW (0.92 g, 4.8 mmol) was dissolved in methanol (100 mL) at 0°C and a stream of
20 hydrogen chloride gas was bubbled into the solution for 10 minutes. The mixture was then stirred for 2 hours at 0°C. The solvent was removed under reduced pressure to give 1 g of the desired compound as a colorless oil.

Analysis Calculated for C₁₀H₁₁N₃O₂:

25 C, 58.53; H, 5.40; N, 20.48.
Found: C, 58.27; H, 5.35, N, 20.13.

Example AY

30



The compound of Example AX (800 mg, 3.9 mmol) was dissolved in EtOH (30 mL) and transferred to a Parr Shaker

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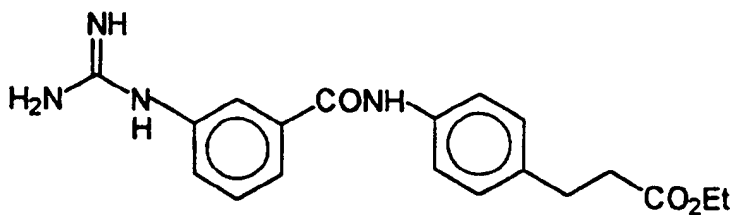
- with 4% Pd/C (200 mg). The reaction was shaken for 24 hours at room temperature under 60 psi pressure of H₂. The reaction mixture was filtered and concentrated and the residue was dissolved in 4 N HCl dioxane solution (4 mL).
- 5 The solvent was removed and the residue was recrystallized from ether to give 550 mg of pure title compound as white solid.

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Example 63

Synthesis of ethyl 4-[[[3-[(aminoiminomethyl)amino]
phenyl]carbonyl]amino]benzenepropanoate, monohydrate
5 trifluoroacetate salt

10



15

The title compound was prepared in the same manner as
described in Example 34, replacing the compound of Example
AA with the compound of Example F.

Analysis Calculated for $C_{19}H_{22}N_4O_3 \cdot 1.1 \text{ TFA} \cdot 1 \text{ H}_2\text{O}$:

C, 51.15; H, 5.08; N, 11.25.

Found: C, 51.17; H, 4.54; N, 11.40.

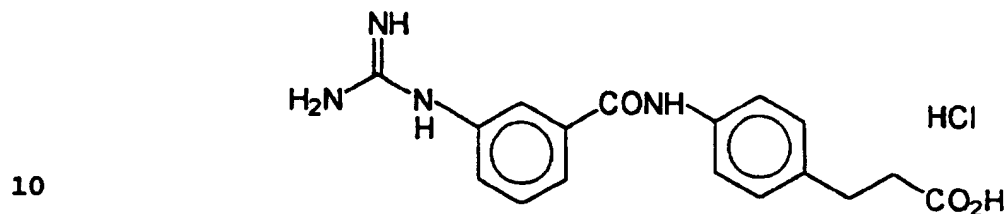
20

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Example 64

Synthesis of 4-[[[3-[(aminoiminomethyl)amino]phenyl]
carbonyl]amino]benzenepropanoic acid, hydrochloride

5



The compound of Example 63 was hydrolyzed in the same manner as described in Example 62.

15 Analysis Calculated for C₁₇H₁₈N₄O₃·0.4 H₂O·1 HCl:

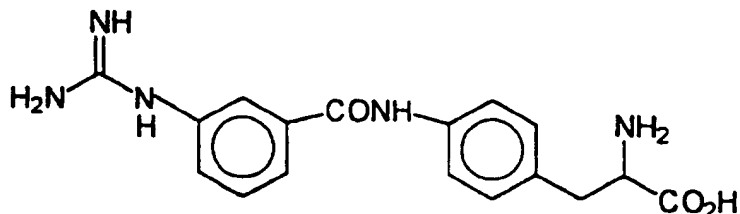
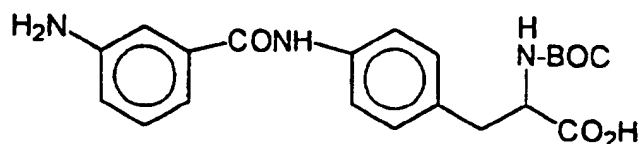
C, 55.18; H, 5.39; N, 15.14.

Found: C, 55.25; H, 5.35; N, 15.00.

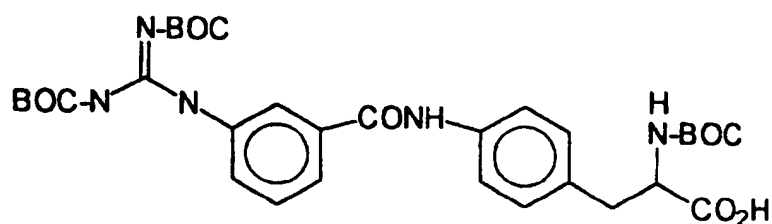
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Example 65

Synthesis of 4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]amino]phenylalanine, trifluoroacetate salt

Step A

The compound of Example AJ was coupled to 3-nitrobenzoyl chloride as described in Example AQ. The resulting nitro compound was hydrogenated in the same manner as described in Example AA to produce the above compound.

Step B

The above compound was synthesized under conditions similar to those described in Example AS using the product of Step A.

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Step C

To a solution of the product of Step B (220 mg, 0.34 mmol) in CH_2Cl_2 (5 mL) was added trifluoroacetic acid (1 mL). The reaction was stirred for 4 hours, concentrated
5 and purified by reverse phase HPLC to afford the title compound.

Analysis Calculated for $\text{C}_{17}\text{H}_{19}\text{N}_5\text{O}_3 \cdot 2.1 \text{ TFA} \cdot 0.2 \text{ H}_2\text{O}$:

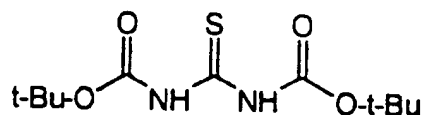
C, 43.57; H, 3.71; N, 11.98.

10 Found: C, 43.26; H, 3.49; N, 11.75.

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Example AZ

5

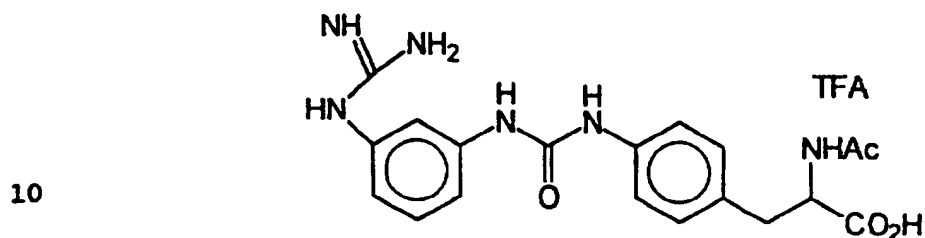
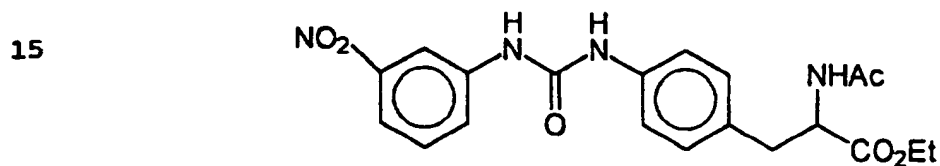


An oven-dried flask equipped with a stirring bar was charged with NaH (60% in mineral oil, 3.36 g, 140 mmol). The NaH was trituated with hexane followed by THF (distilled) under argon. The NaH was then suspended in THF (250 mL) and the mixture was cooled to 0°C. Thiourea (1.426 g, 18.7 mmol) was added in one portion. After 5 minutes, the reaction was allowed to stir at room temperature for 10 minutes. The reaction was cooled to 0°C and di-*t*-butyl dicarbonate (9.001 g, 41.2 mmol) was added. Within 15 minutes of addition, the reaction became a tan slurry. The reaction was allowed to warm slowly to room temperature over 2 hours and then stirred at room temperature for an additional 1 hour. The reaction was quenched with saturated NaHCO₃ (50 mL) and poured into water (600 mL). The mixture was extracted with EtOAc (3X150 mL) and the organic layers were combined and dried over Na₂SO₄. Concentration *in vacuo* gave the product as a yellow solid (4.23 g, 82% yield). NMR was consistent with the proposed structure.

- 150 -

Example 68

Synthesis of N-acetyl-4-[[[3-[(aminoiminomethyl)amino]phenyl]amino]carbonyl]amino]phenylalanine,
5 trifluoroacetate salt

Step A

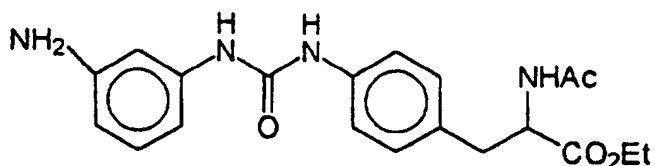
To a stirred solution of 3-nitrophenylisocyanate
20 (0.082 g, 0.5 mmol, Aldrich) in methylene chloride (5 mL) was added to the product from Example AA (0.1 g, 0.4 mmol) in small portions over 5 minutes. The mixture was stirred 18 hours at room temperature. The mixture was then poured into 10% aqueous sodium
25 hydroxide (50 mL) and washed with ethyl acetate (2 x 25 mL). The basic solution was acidified with 10% HCl and the resulting precipitate was filtered and dried. This produced 0.16 g (97%) of the title compound.

30 HRMS (MH⁺) for C₂₀H₂₃N₄O₆ Calculated: 415.1618
Found: 415.1654

- 151 -

Step B

5



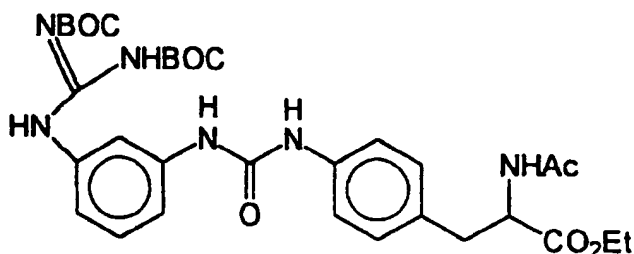
A stirred solution of the product of Example 68A (0.16 g, 0.39 mmol) in ethyl alcohol (25 mL) and THF (50 mL) was hydrogenated over 4% palladium on carbon under an atmosphere of hydrogen at 5 psi. The solvent was removed at reduced pressure. This produced 0.14 g (93%) of the title compound.

HRMS (M+) for $C_{20}H_{24}N_4O_4$	Calculated:	384.1797
	Found:	384.1837

15

Step C

20

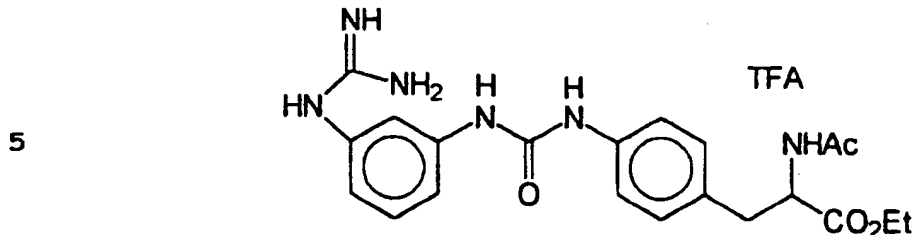


The product from Example 68B (0.14 g, 0.36 mmol) was subjected to the reaction conditions described for the preparation of Example AS. The crude product was chromatographed on silica gel eluting with ethyl acetate which produced 0.21 g (93%) of the title compound.

30

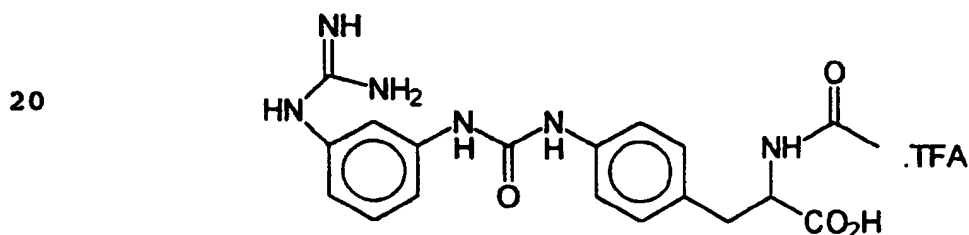
APCI MS (MH+) for $C_{31}H_{43}N_6O_8$	Calculated:	627
	Found:	627

- 152 -

Step D

10 The product from Example 68C (0.21 g, 0.34 mmol) was subjected to the reaction conditions described for the preparation of Example 60. This produced 0.17 g (93%) of the title compound.

15 APCI MS (free base MH+) for $C_{21}H_{27}N_6O_4$ Calculated: 427
Found: 427

Step E

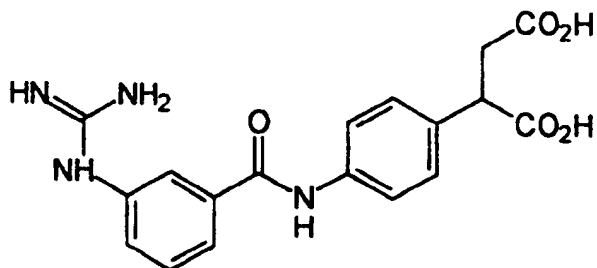
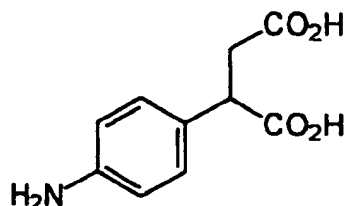
25 The product of Example 68D (0.17 g, .31 mmol) in methyl alcohol (1 mL) was cooled (0°C) and treated with 1 N lithium hydroxide solution (0.7 mL, 0.7 mmol). The solution was warmed to room temperature and stirred 18 hours. The volatile components were removed at reduced
30 pressure on a rotary evaporator. The crude product was chromatographed (reverse phase HPLC, gradient elution with water/acetonitrile/trifluoroacetic acid). This produced 100 mg (63%) of the title compound.

35 ESI MS (free base MH+) for $C_{19}H_{23}N_6O_4$ Calculated: 399
Found: 399

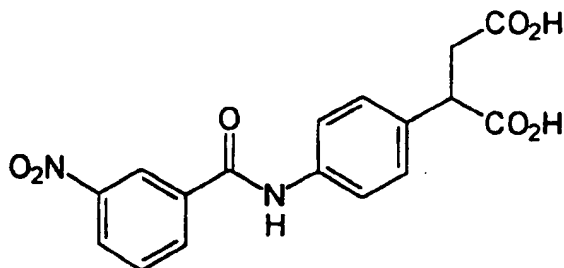
- 153 -

Example 69

Synthesis of [4-[[[3-[(aminoiminomethyl)amino]-
phenyl]carbonyl]amino]phenyl]butanedioic
acid, trifluoroacetate salt

Step A

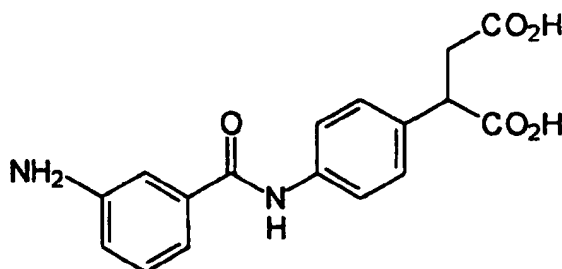
4-Nitrophenylsuccinic acid (Lancaster) (5 g, 2.0
mmol) was added to absolute ethanol (70 mL) in a Parr
jar. Palladium on carbon 5% (700 mg) was added and the
mixture was hydrogenated under 50 psi in a Parr
apparatus for a period of 2.5 hours. After complete
reaction the palladium catalyst was removed under
reduced pressure and the sample dried in vacuo to give
a white colored solid (5 g, 99% yield). NMR and MS
were consistent with the proposed structure.

Step B

- 154 -

The 4-anilinophenylsuccinic acid (2 g) from Step A was placed in a flask followed by water (50 mL) and KOH (1 g). To this solution 3-nitrobenzoyl chloride (Aldrich) dissolved in acetonitrile (10 mL) was added dropwise. The reaction was monitored by HPLC. After the reaction was complete (1 hour), 10% HCl was added and the solution was extracted with ethyl acetate and dried over Na_2SO_4 . After evaporation of the solvent a white solid remained (1.5 g). NMR and MS were consistent with the proposed structure.

Step C



20

The white solid from Step B (1.5 g) was added to absolute ethanol (50 mL) and acetic acid (2 mL) in a Parr jar. Palladium on carbon 5% (500 mg) was added and the mixture was hydrogenated under 50 psi in a Parr apparatus for a period of 1.5 hours. After complete reaction the palladium catalyst was removed by filtration through a plug of celite. The solvent was removed under reduced pressure and the sample dried in vacuo to give a white colored solid (1.5 g, 99% yield). NMR and MS were consistent with the proposed structure.

Step D

A portion of the white solid from Step C above (400 mg) was added to acetonitrile (20 mL) followed by pyrazole carboxamidine HCl (Aldrich) (1 g) and DIEA. The mixture was heated to reflux for 4 hours. After the reaction was complete water was added and TFA was

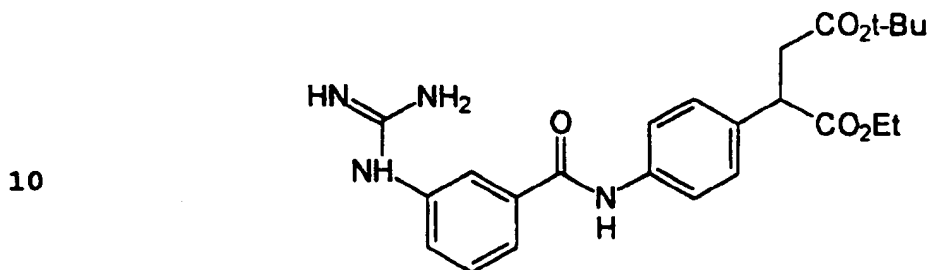
- 155 -

added to bring the pH=2. The product was purified by reverse phase chromatography (water/acetonitrile) to result in the title compound as a white solid (320 mg). NMR and MS were consistent with the proposed structure.

- 156 -

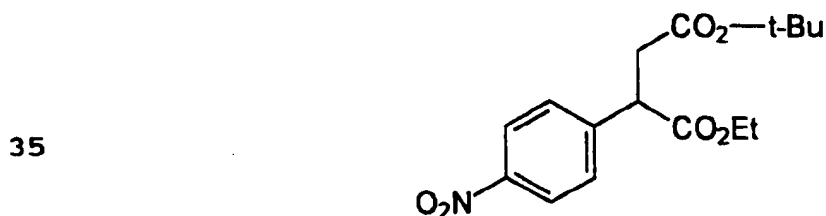
Example 70

Synthesis of 4-(1,1-dimethylethyl) 1-ethyl[4-[[[3-
[(aminoiminomethyl)amino]phenyl]carbonyl]amino]phenyl]-
5 butanedioate, trifluoroacetate salt

Step A

4-Nitrophenylacetic acid (Aldrich) (10 g) was
20 added to a flask containing ethanol (200 mL) and acetyl
chloride (20 mL). The solution was heated to reflux
for 1 hour then left to stir overnight at room
temperature. The solvent was removed under reduced
pressure to result in a solid which was dissolved in
25 hexane (150 mL) and washed with water (100 mL). The
hexane solution was dried over Na₂SO₄ and the solvent
removed under reduced pressure to give the ethyl ester
(12 g) as a white solid. NMR and MS were consistent
with the proposed structure.

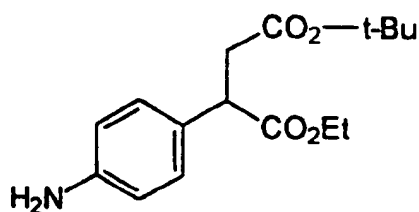
30

Step B

- 157 -

Ethyl 4-nitrophenylacetate (5 g) from Step A was dissolved in THF (100 mL). Potassium tert-butoxide (45 mL) was added at 0°C. The solution turned a deep purple indicating the anion was formed. After 15 minutes at 0°C tert-butyl bromacetate (5.2 g) was added. After the reaction was complete 10% HCl was added and the product extracted with ethyl acetate, and dried over Na₂SO₄ to give a dark yellow oil (7 g). NMR and MS were consistent with the proposed structure.

Step C

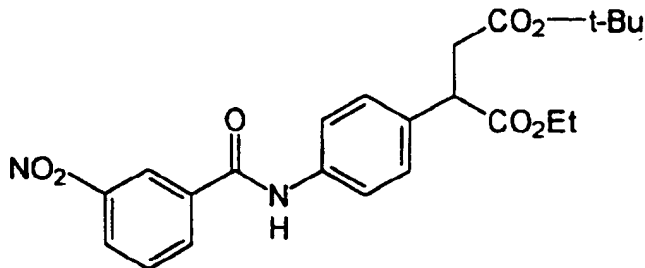


The compound from Step B (5 g) was added to absolute ethanol (70 mL) and acetic acid (2 mL) in a Parr jar. Palladium on carbon 5% (700 mg) was added and the mixture was hydrogenated under 50 psi in a Parr apparatus for a period of 2.5 hours. After complete reaction the palladium catalyst was removed by filtration through a plug of celite. The solvent was removed under reduced pressure and the sample dried in vacuo to give a white colored solid (5 g, 99% yield). NMR and MS were consistent with the proposed structure.

- 158 -

Step D

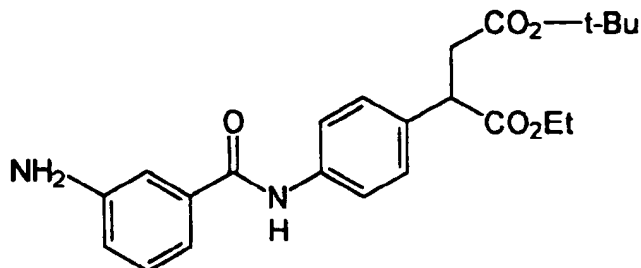
5



10 The 4-anilinophenylsuccinic diester (2 g) from
Step C was placed in a flask followed by water (50 mL)
and K_2CO_3 (1 g). To this solution 3-nitrobenzoyl
chloride (Aldrich) dissolved in acetonitrile (10 mL)
was added dropwise. The reaction was monitored by
15 HPLC. After the reaction was complete (1 hour), 10%
HCl was added and a white solid was separated (1.5 g).
NMR and MS were consistent with the proposed structure.

Step E

20



25

 The compound from Step D (1 g) was added to
absolute ethanol (70 mL) and acetic acid (2 mL) in a
30 Parr jar. Palladium on carbon 5% (700 mg) was added
and the mixture was hydrogenated under 50 psi in a Parr
apparatus for a period of 2.5 hours. After complete
reaction the palladium catalyst was removed by
filtration through a plug of celite. The solvent was
35 removed under reduced pressure and the sample dried in
vacuo to give a white colored solid (1 g, 99% yield).
NMR and MS were consistent with the proposed structure.

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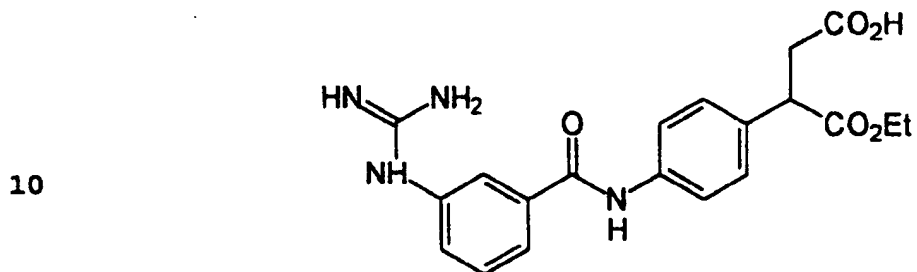
Step F

A portion of the white solid from Step E (1 g) was added to acetonitrile (20 mL) followed by pyrazole carboxamidine HCl (Aldrich) (2 g) and DIEA (2 g). The mixture was heated to reflux for 6 hours. After the reaction was complete water was added and TFA was added to bring the pH=2. The product was purified by reverse phase chromatography (water/acetonitrile) to result in the title compound as a white solid (820 mg). NMR and MS were consistent with the proposed structure.

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Example 71

Synthesis of 1-ethyl hydrogen[4-[[[3-
[(aminoiminomethyl)amino]phenyl]carbonyl]amino]-
5 phenyl]butanedioate, trifluoroacetate salt

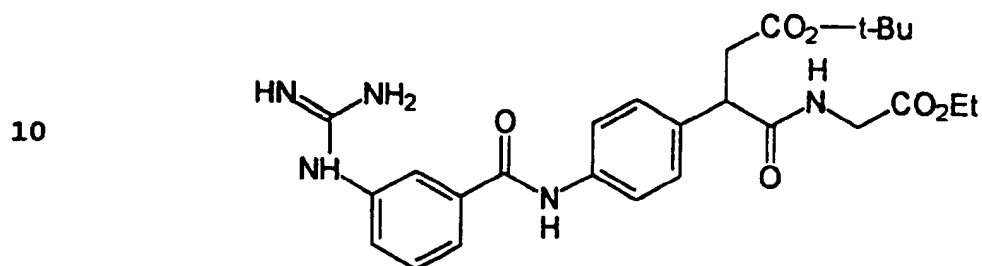
Step A

The compound from Step D in Example 70 (500 mg)
15 was added to methylene chloride (5 mL) followed by TFA
(2 mL). The reaction was monitored by HPLC. After the
reaction was complete the product was purified by
reverse phase chromatography (water/acetonitrile) to
result in a white solid (310 mg). NMR and MS were
20 consistent with the proposed structure.

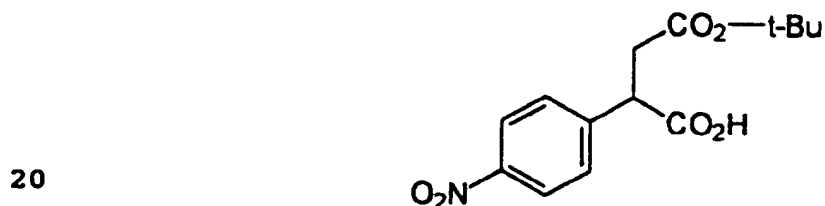
- 161 -

Example 72

Synthesis of 1,1-dimethylethyl 4-[[[3-
[(aminoiminomethyl)amino]phenyl]carbonyl]-
5 amino]- β -[[[(ethoxycarbonyl)methyl]amino]-
carbonyl]benzenepropanoate, trifluoroacetate salt

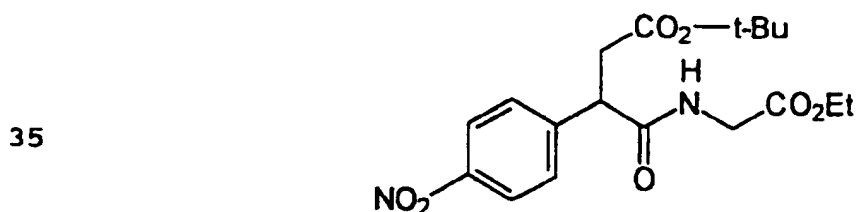


15 Step A



The product from Step B (2 g) of Example 70 was added to water/acetonitrile 1:1 (50 mL), followed by the addition of lithium hydroxide (100 mg, 0.4 mmol).
25 The reaction was stirred at 25°C, and monitored by HPLC. After complete hydrolysis the product was extracted with ethyl acetate dried over Na₂SO₄ to give a yellow oil (1.7 g). NMR and MS were consistent with the proposed structure.

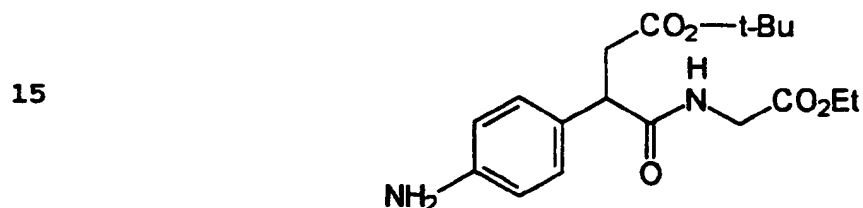
30

Step B

- 162 -

N,N'-Disuccinimidyl carbonate (1.7 g) was added to the 4-nitro-phenyl adduct from Step A (1.5 g) in dry dimethylformamide (20 mL) followed by dimethylaminopyridine (200 mg). After a period of 1 hour glycine ethyl ester hydrochloride (1 g) in DMF (10 mL) and NMM (2 mL) was added in one portion. After complete reaction, the product was extracted into ethyl acetate and worked up to give the desired product (1.5 g). NMR and MS were consistent with proposed structure.

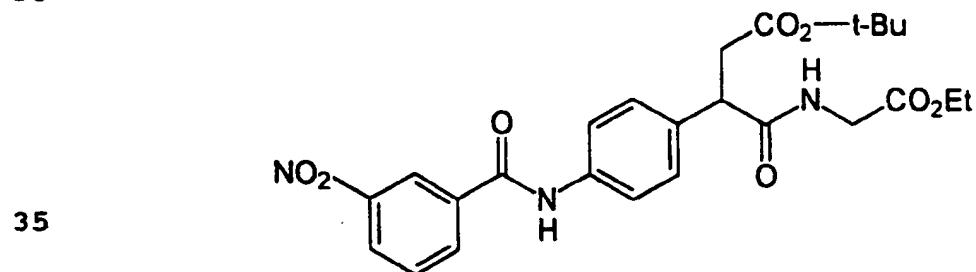
Step C



The compound from Step B (1 g) was added to absolute ethanol (70 mL) in a Parr jar. Palladium on carbon 5% (500 mg) was added and the mixture was hydrogenated under 50 psi in a Parr apparatus for a period of 2.5 hours. After complete reaction the palladium catalyst was removed by filtration through a plug of celite. The solvent was removed under reduced pressure and the sample dried in vacuo to give a white colored solid (1 g, 99% yield). NMR and MS were consistent with the proposed structure.

Step D

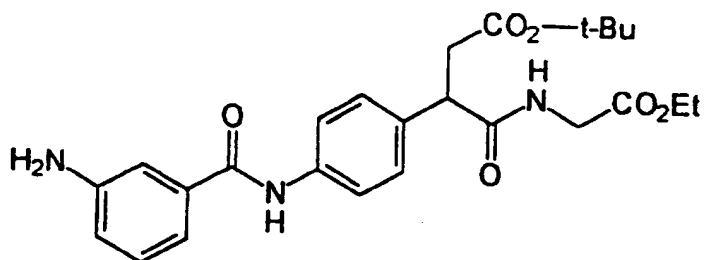
30



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The product from Step C (1 g) was placed in a flask, followed by addition of water (50 mL) and K_2CO_3 (1 g). To this solution 3-nitrobenzoyl chloride (Aldrich) dissolved in acetonitrile (10 mL) was added dropwise. The reaction was monitored by HPLC. After the reaction was complete (1 hour), 10% HCl was added and the solution was extracted with ethyl acetate and dried over Na_2SO_4 . After evaporation of the solvent a white solid remained (1.56 g). NMR and MS were consistent with the structure.

Step E



The compound from Step D (1.2 g) was added to absolute ethanol (70 mL) and acetic acid (2 mL) in a Parr jar. Palladium on carbon 5% (700 mg) was added and the mixture was hydrogenated under 50 psi in a Parr apparatus for a period of 2.5 hours. After complete reaction the palladium catalyst was removed by filtration through a plug of celite. The solvent was removed under reduced pressure and the sample dried in vacuo to give a white colored solid (1.1 g, 99% yield). NMR and MS were consistent with the proposed structure.

Step F

A portion of the white solid from Step E (1 g) was added to acetonitrile (20 mL) followed by pyrazole carboxamidine HCl (Aldrich) (2 g) and DIEA. The mixture was heated to reflux for 6 hours. After the reaction was complete water was added and TFA was added

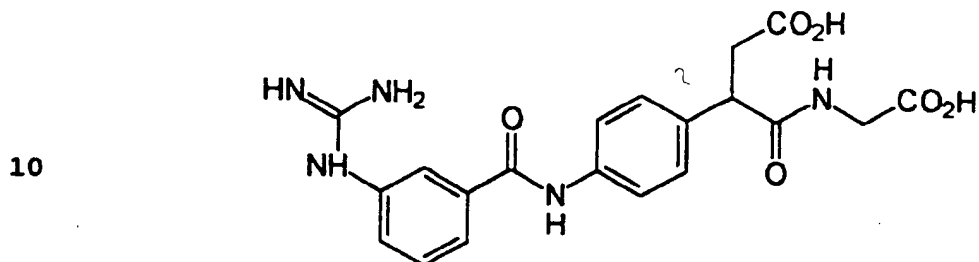
- 164 -

to bring the pH=2. The product was purified by reverse phase chromatography (water/acetonitrile) to result in the title compound as a white solid (700 mg). NMR and MS were consistent with the structure.

- 165 -

Example 73

Synthesis of 4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]amino]-beta-[[[(carboxymethyl)amino]carbonyl]-
5 benzenepropanoic acid, trifluoroacetate salt

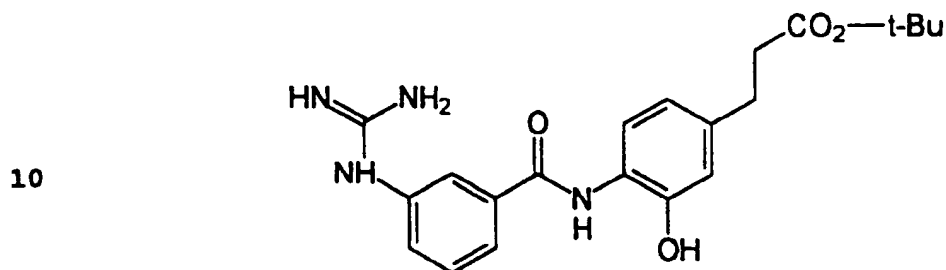
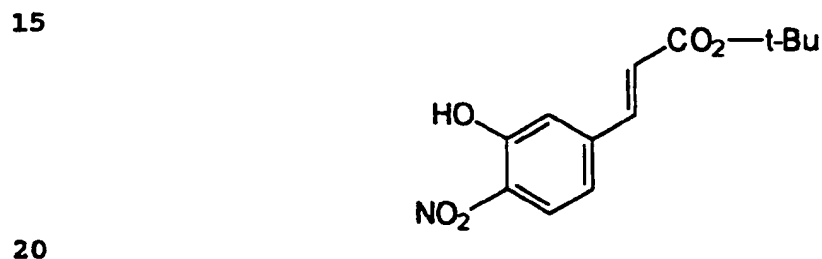
Step A

15 The compound of Example 72 (500 mg) was added to
methylene chloride (5 mL) followed by the addition of
TFA (2 mL). The reaction was monitored by HPLC. After
the reaction was complete the product was freeze dried.
The product was added to water/aceonitrile 1:1 (50 mL),
20 followed by the addition of lithium hydroxide (100 mg,
0.4 mmol). The reaction was stirred at 25°C, and
monitored by HPLC. After complete hydrolysis the
product was extracted with ethyl acetate and dried over
Na₂SO₄ to give a yellow oil (450 mg). NMR and MS were
25 consistent with the proposed structure.

- 166 -

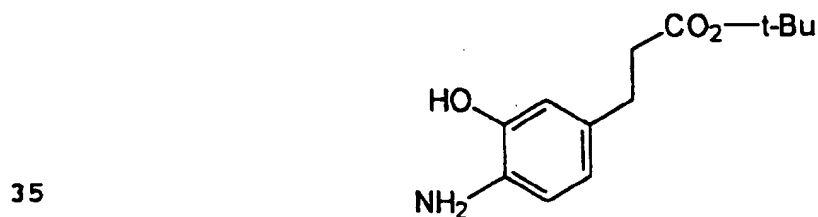
Example 74

Synthesis of 1,1-dimethylethyl 4-[[[3-
[(aminoiminomethyl)amino]phenyl]carbonyl]amino]-3-
5 hydroxybenzenepropanoate, trifluoroacetate salt

Step A

To 4-nitro-3-hydroxy benzaldehyde (Aldrich) (5 g)
in acetonitrile (40 mL) was added (tert-
butoxycarbonylmethylene)triphenylphosphorane (Aldrich)
25 (11 g) and DBU (1 mL). The reaction was stirred for 1
hour. The solvent was concentrated to 20 mL and the
product crystallized from the solution (4 g). NMR and
MS were consistent with the proposed structure.

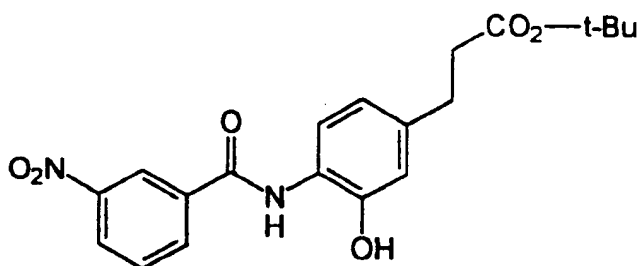
30 Step B



- 167 -

The compound from Step A (3 g) was added to absolute ethanol (70 mL) in a Parr jar. Palladium on carbon 5% (700 mg) was added and the mixture was hydrogenated under 50 psi in a Parr apparatus for a period of 2.5 hours. After complete reaction the palladium catalyst was removed by filtration through a plug of celite. The solvent was removed under reduced pressure and the sample dried in vacuo to give a white colored solid (3 g, 99% yield). NMR and MS were consistent with the proposed structure.

Step C

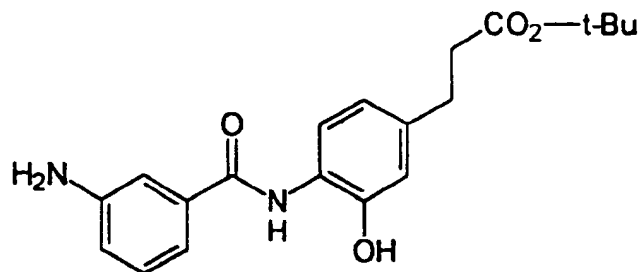


The product from Step B (1.5 g) was placed in a flask followed by the addition of methylenechloride (50 mL) and NMM (2 g). To this solution 3-nitrobenzoylchloride (Aldrich) was added. The reaction was monitored by HPLC. After the reaction was complete (1 hour) 10% HCl was added and the solution was extracted with ethyl acetate and dried over Na_2SO_4 . After evaporation of the solvent a white solid remained (3 g). NMR and MS were consistent with the proposed structure.

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Step D

5



10 The compound from Step C (1 g) was added to absolute ethanol (70 mL) and acetic acid (2 mL) in a Parr jar. Palladium on carbon 5% (500 mg) was added and the mixture was hydrogenated under 50 psi in a Parr apparatus for a period of 2.5 hours. After complete
15 reaction the palladium catalyst was removed by filtration through a plug of celite. The solvent was removed under reduced pressure and the sample dried in vacuo to give a white colored solid (1 g, 99% yield). NMR and MS were consistent with the proposed structure.

20

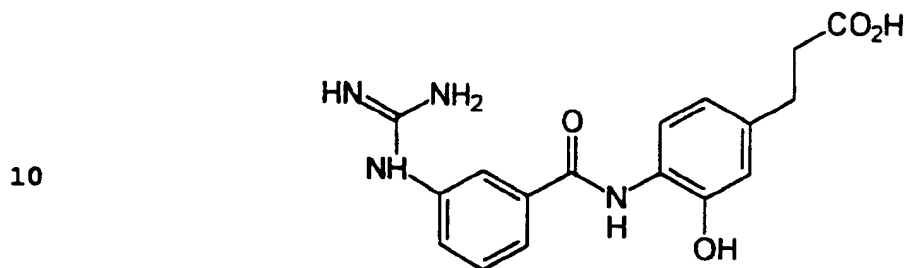
Step E

 The product from Step D (1 g) was added to acetonitrile (20 mL) followed by pyrazole carboxamidine HCl (Aldrich) (23 g) and DIEA. The mixture was heated
25 to reflux for 6 hours. After the reaction was complete, water and TFA were added to bring the pH=2. The product was purified by reverse phase chromatography (water/acetonitrile) to result in the title compound as a white solid (700 mg). NMR and MS
30 were consistent with the proposed structure.

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Example 75

Synthesis of 4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]amino]-3-hydroxybenzenepropanoic acid,
5 trifluoroacetate salt

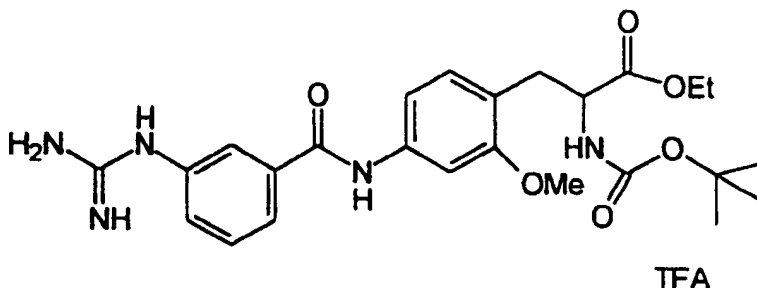
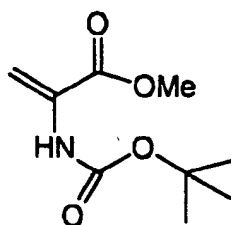
Step A

The compound of Example 74 (300 mg) was added to
15 methylene chloride (5 mL) followed by the addition of
TFA (2 mL). The reaction was monitored by HPLC. After
the reaction was complete (2 hours) the product was
purified by reverse phase chromatography
(water/acetonitrile) to result in a white solid (110
20 mg). NMR and MS were consistent with the proposed
structure.

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Example 76

4-[[[3-[(Aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(1,1-dimethylethoxy)carbonyl]-
2-methoxyphenylalanine, ethyl ester

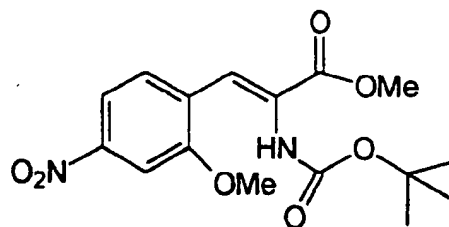
Step A

To a solution of methyl pyruvate (2.042 g, 20 mmol) in benzene (70 mL) was added tert-butyl carbamate (9.372 g, 80 mmol) and the resulting suspension was heated to obtain full dissolution. The mixture was heated to 70°C and phosphorus oxychloride (7.3 mL, 80 mmol) was added dropwise over 1 minute. After 2 minutes a heavy white precipitate appeared and the reaction mixture was stirred at 70°C for an additional 15 minutes. The mixture was rapidly cooled, poured into an aqueous solution of sodium dihydrogenphosphate (pH 4) and the aqueous layer extracted three times with dichloromethane. The organic layer was washed with water, dried and evaporated under reduced pressure. Chromatography of the resulting oil on silica gel eluting with dichloromethane afforded 1.6 g of the product.

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Step B

5



To 5 mL of dimethylformamide was added the
10 compound of Step A (600 mg, 2.98 mmol), 2-bromo-5-nitroanisole (494 mg, 2.13 mmol), tetrabutylammonium chloride (591 mg, 2.13 mmol), sodium bicarbonate (447 mg, 5.3 mmol) and 2 mg of palladium(II) acetate. The slurry was degassed with argon, sealed and heated at
15 85°C overnight. The reaction mixture was diluted with water and extracted with dichloromethane. The combined extracts were washed once with water, dried (MgSO₄) and concentrated to give the crude product. Chromatography on silica gel using 20/80 ethyl acetate/hexane as
20 eluent gave 600 mg of product.

Analysis Calculated for C₁₆H₁₉N₂O₇:

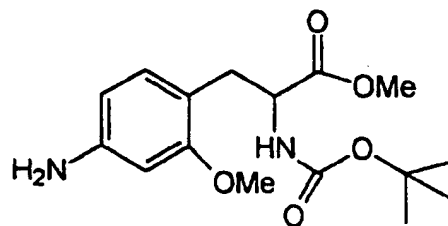
C, 54.70; H, 5.45; N, 7.97.

Found: C, 54.43; H, 5.64; N, 8.09.

25

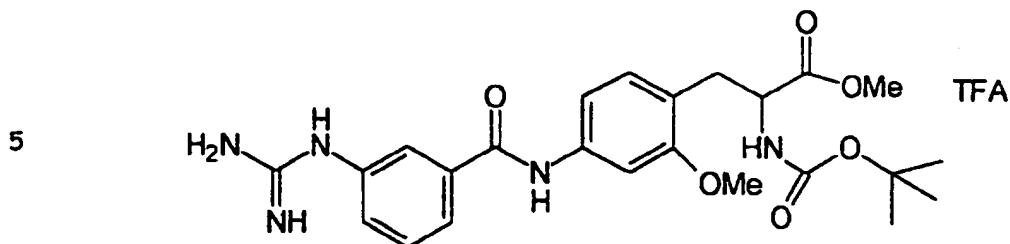
Step C

30



The compound of Step B was shaken in a Parr
35 apparatus in ethanol with 5% Pd on carbon at room temperature under 5 psi H₂ for 16 hours. The solution was filtered and concentrated to give the product.

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Step D

To 10 mL of dimethylformamide was added 3-
10 guanidinobenzoic acid HCl (Example A) (299 mg, 1.387 mmol) and N-methylpiperidine (138 mg, 1.387 mmol). The reaction mixture was cooled to 0°C and isobutylchloroformate (189 mg, 1.387 mmol) was added. After 5 minutes the compound of Step C (450 mg, 1.387
15 mmol) was added and the reaction mixture was stirred at room temperature overnight. The reaction mixture was concentrated and purified by HPLC on a C18 column using a MeCN/H₂O gradient as eluant to give a white solid (495 mg).

20

Analysis Calculated for C₂₄H₃₁N₅O₆+TFA+0.3 H₂O:

C, 51.62; H, 5.43; N, 11.58.

Found: C, 51.27; H, 5.27; N, 11.49.

25 Step E

To the compound of Step D in 3 ml of methanol was added 1.0 mL of 1M LiOH. The reaction mixture was stirred at room temperature overnight. The solvent was removed under vacuum and 10 mL of water was added. The
30 aqueous layer was acidified with trifluoroacetic acid and purified by HPLC on a C18 column using a MeCN/H₂O gradient as eluant to give a white solid (89 mg).

35

Analysis Calculated for C₂₃H₂₉N₅O₆+TFA+0.6 H₂O:

C, 50.35; H, 5.27; N, 11.74.

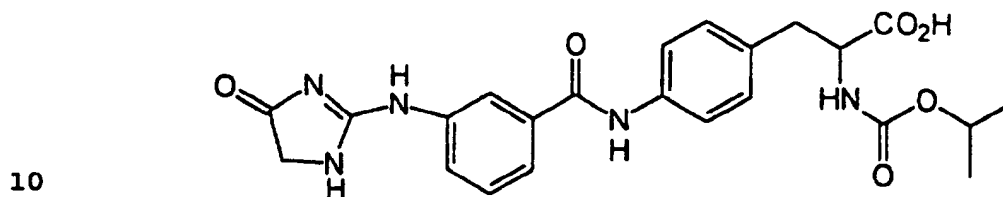
Found: C, 50.23; H, 5.13; N, 11.71.

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Example 77

4-[[[3-[(4,5-dihydro-4-oxo-1H-imidazol-2-yl)amino]-
phenyl]carbonyl]amino]-N-[(1-methylethoxy)-
carbonyl]phenylalanine

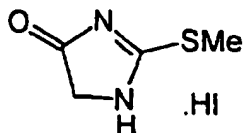
5



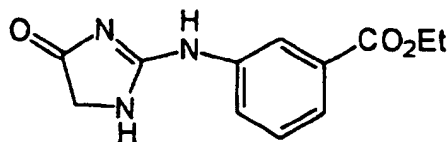
10

Step A

15



To a mixture of 2-thiohydantoin (5.5 g, 47.4 mmol) in absolute ethanol (60 mL) was added methyl iodide (3.5 mL, 56.6 mmol). The mixture was heated at reflux for 5 hours. The mixture was cooled to room temperature and concentrated *in vacuo* and the crude product used directly in the next step.

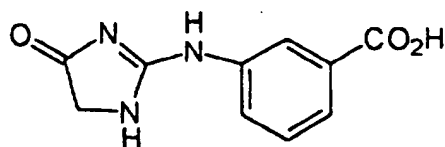
25 Step B

30

To a mixture of the thiomethyl compound from Step A (1.0 g, 3.8 mmol) in absolute ethanol (20 mL) was added ethyl 3-aminobenzoate (2.5 g, 15.3 mmol). The mixture was stirred at room temperature for 16 hours. The mixture was concentrated *in vacuo* and the residue chromatographed (85:14:1 CH₂Cl₂:MeOH:NH₄OH) to give the desired product (414 mg, 44%).

35

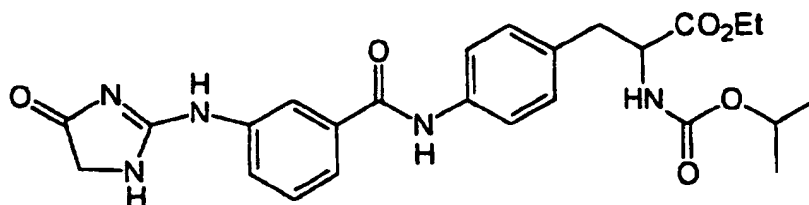
- 174 -

Step C

5

To a mixture of the ester (250 mg, 1.0 mmol) in THF (2 mL) and methanol (2 mL) was added 1.0 N NaOH solution (2 mL). The reaction solution was stirred at room temperature for 2 hours and concentrated *in vacuo*. The residue was suspended in water and carefully acidified to pH 4 with 1 N HCl. The solid was collected by filtration and washed with water and ether to give the desired product (190 mg, 87%).

15

Step D

20

To a solution of the acid from Step C (150 mg, 0.7 mmol) in DMF (4 mL) at 0°C was added 1-methylpiperidine (15 mL, 1.4 mmol) and isobutylchloroformate (0.088 mL, 0.7 mmol). The solution was stirred for 5 minutes and the amine from Example AE (186 mg, 0.7 mmol) was added. The mixture was stirred for 20 hours and then concentrated *in vacuo*. The residue was chromatographed (85:14:1 CH₂Cl₂:MeOH:NH₄OH) to give the desired product (73 mg, 22%).

30

Analysis Calculated for C₂₅H₂₉N₅O₆ + 0.25 H₂O:

C, 60.05; H, 5.94; N, 14.00.

35

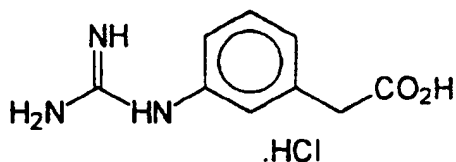
Found: C, 60.10; H, 6.11; N, 14.11.

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Step E

To a mixture of the ester from Step D (40 mg, 0.08 mmol) in THF (0.5 mL) and methanol (0.5 mL) was added
5 1.0 N NaOH solution (0.4 mL). The solution was stirred at room temperature for 3 hours and concentrated *in vacuo*. The residue was dissolved in water and carefully acidified to pH 4 with 1 N HCl. The white solid was collected by filtration to give the desired
10 compound (27 mg, 71%). ¹H NMR (CD₃OD) δ 7.19-7.90 (m, 8H), 4.75 (m, 1H); 4.34 (m, 1H); 3.95 (s, 2H); 3.13 (m, 1H); 2.91 (m, 1H); 1.18 (d, 3H, J = 7.2 Hz); 1.12 (d, 3H, J = 7.2 Hz).

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Example BJ

5

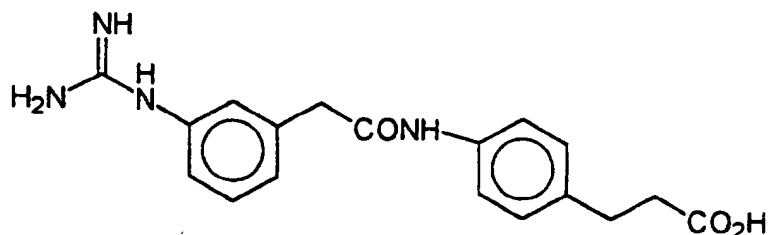
A solution of 3-aminophenylacetic acid (2.712 g, 17.9 mmol), Example AM (3.023 g, 20.6 mmol), and Hunig's base (3.6 mL, 20.6 mmol) in dioxane (30 mL)/water (15 mL) was refluxed for 16 hours under argon. With heating, a white precipitate was observed. The reaction was cooled to room temperature and the white solid filtered off. The solid was washed with 1:1 dioxane/water (3 X 5 mL). The solid was suspended in 15 mL of water and acidified with concentrated HCl until the solid dissolved. The solution was concentrated in vacuo and the resulting yellow residue slurried with ether. The yellow solid was collected by vacuum filtration (3.025 g, 74% yield). NMR was consistent with proposed structure.

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Example 78

Synthesis of 4-[[2-[4-[(aminoiminomethyl)amino]phenyl]-
acetyl]amino]benzenepropanoic acid, monohydrate
5 trifluoroacetate salt

10



15

The compound of Example F (497 mg, 1.66 mmol) was
coupled with the compound of Example BJ (380 mg, 1.66
mmol) according to the procedure described in Example
34. The crude product was purified by HPLC-Method 1 to
give the desired product. The product was hydrolyzed
under the conditions described in Example 35 and
purified by HPLC-Method 1 to give a white solid.

20

Analysis Calculated for $C_{18}H_{20}N_4O_3 \cdot 1.5 \text{ TFA} + 1 \text{ H}_2\text{O}$:

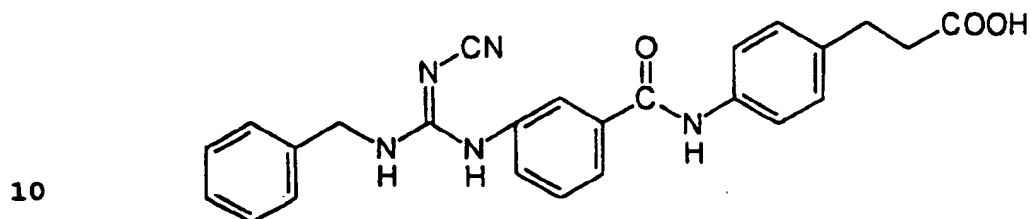
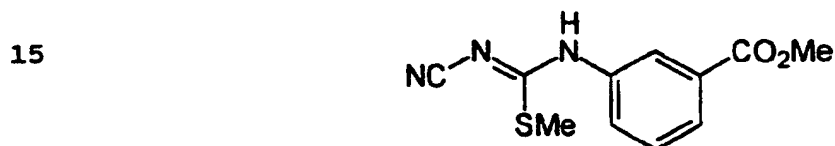
C, 47.64; H, 4.47; N, 10.58.

Found: C, 47.56; H, 4.49; N, 10.77.

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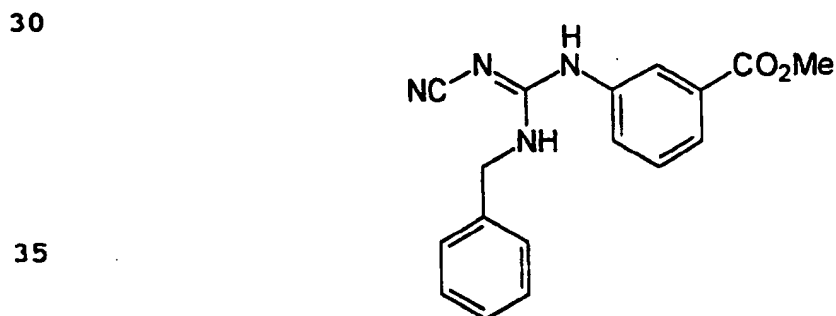
Example 79

Synthesis of 4-[[[3-[[[(cyanoimino)[(phenylmethyl)-
amino]methyl]amino]phenyl]carbonyl]amino]-
5 benzenepropanoic acid

Step A

A stirred mixture of 3-amino methyl benzoate (6.04
20 g, 40 mmol) and dimethyl N-cyanodithioiminocarbonate
(11.96 g, 80 mmol) in pyridine (70 ml) was heated at
reflux under a nitrogen atmosphere for 2.5 hours. The
reaction mixture was cooled to room temperature. On
standing overnight at room temperature the title
25 compound crystallized from the reaction mixture
affording 6.2 g (two crops).

NMR was consistent with the proposed structure.

Step B

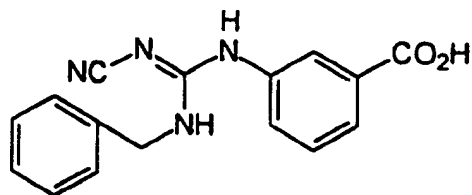
- 179 -

A stirred mixture of the compound from Step A (1.0 g) and benzylamine (440 mg) in ethanol (15 ml) was heated at reflux under a nitrogen atmosphere for 3 hours. The reaction mixture was cooled to room temperature. On standing overnight at room temperature a white solid was obtained and isolated by filtration (720 mg). The crude filtrate was further purified by chromatography on silica (eluant; ethyl acetate/hexane, 1:1) to afford the above compound (550 mg) as a white solid.

NMR was consistent with the proposed structure.

Step C

15



To a stirred solution of the compound from Step B (250 mg) in THF (2 ml) and MeOH (2 ml), 1N-NaOH (2 ml) was added. The reaction mixture was stirred at room temperature for 2 hours and concentrated *in vacuo* to afford a white solid. The residue was acidified by suspension in water followed by addition of 1N-HCl. The resultant solid was filtered, washed with diethyl ether and dried to afford the above compound (140 mg) which was used in the next step without further purification.

30

NMR was consistent with the proposed structure.

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Step D

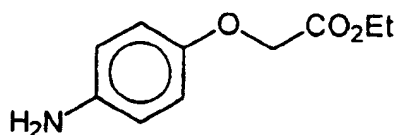
To a stirred solution of the product of Step C (440 mg, 1.5 mmol), in methylene chloride (20 mL) at 0°C, triethylamine (0.8 mL), DMAP (20 mg), EDCI (288 mg, 1.5 mmol) and the compound of Example F (448 mg, 1.5 mmol) were added. The reaction mixture was stirred at 0°C for 15 minutes, allowed to attain room temperature and then stirred for another 16 hours. The reaction mixture was concentrated and the residue was dissolved in EtOAc. Washed with water, saturated aqueous NaHCO₃ and brine, dried over Na₂SO₄ and evaporated. The residue was chromatographed on silica gel using 2:1 Hexane/EtOAc as eluant to give 400 mg of pure ester as white solid. The white solid was dissolved in MeOH (4 mL) and lithium hydroxide (1 M, 2 mL) was added. The reaction mixture was stirred at room temperature overnight, and then concentrated. The residue was dissolved in water and acidified with TFA. White precipitate resulted was filtered, and washed with water and ether to give the title compound as white solid (300 mg).

Analysis Calculated for C₂₅H₂₃N₅O₃·0.1 H₂O:

C, 67.74; H, 5.27; N, 15.80.

Found: C, 67.58; H, 5.09; N, 15.95.

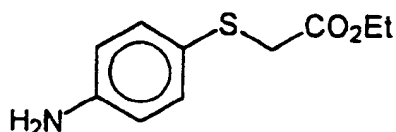
- 181 -

Example BA

5

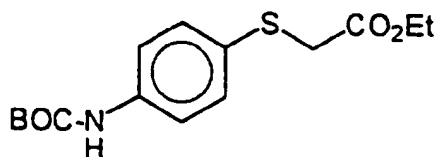
A solution of 4-aminophenol in THF (50 mL) was cooled to -30°C. Sodium hydride was added and the reaction mixture was warmed to 0°C and kept at this temperature for 0.5 hour. The reaction mixture was cooled to -30°C and ethyl bromoacetate was added. The reaction mixture was warmed to room temperature and stirred overnight. The solution was concentrated and partitioned between ethyl ether and water. The organic layer was washed two times with 10% NaOH solution and brine, dried over Na₂SO₄ and evaporated. The crude product was chromatographed on silica gel using CHCl₃/EtOH/NH₄OH (95:5:1) as eluant to give 550 mg of the desired compound as a brown solid.

20

Example BB

25

The above compound was synthesized in the same manner as described in Example BA, replacing 4-aminophenol with 4-aminothiophenol.

Example BC

30

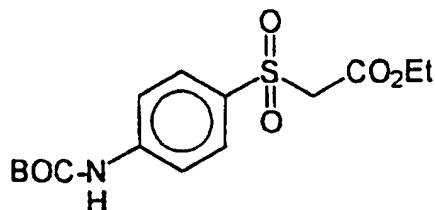
35

The compound of Example BB was protected with t-BOC in the same manner as described in Example X.

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Example BD

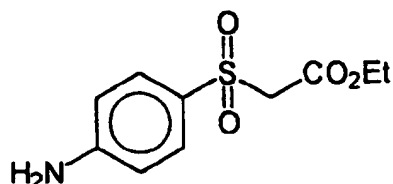
5



The compound of Example BC (0.9 g, 2.3 mmol) was dissolved in methanol (15 mL) and cooled to 0°C. Oxone (potassium peroxomonosulfate) (2.85 g, 4.6 mmol) in water (17 mL) was added and the solution was stirred at room temperature overnight. The solution was diluted with water, extracted with methylene chloride, dried over Na₂SO₄ and concentrated. Crystallization from EtOAc/Hexane resulted in 1.3 g of a white solid.

Example BE

20



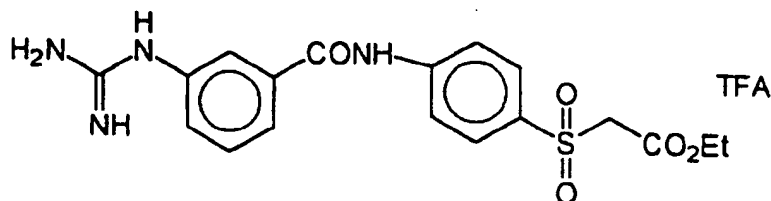
The compound of Example BD (0.8 g, 2.3 mmol) was dissolved in methylene chloride (4 mL) and cooled to 0°C. To the reaction mixture was added trifluoroacetic acid (2 mL). After 15 minutes ice bath was removed and the reaction continuously stirred for 3 hours. The reaction mixture was then concentrated to give 860 mg of desired compound.

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Example 80

Synthesis of ethyl [[4-[[[3-[(aminoiminomethyl)amino]
phenyl]carbonyl]amino]phenyl]sulfonyl]acetate,
5 monohydrate trifluoroacetate salt

10



15

The title compound was prepared in the same manner
as described in Example 34, replacing the compound of
Example AA with the compound of Example BE.

Analysis Calculated for C₁₈H₂₀N₄O₅S · 1 TFA · 1 H₂O:

C, 44.78; H, 4.32; N, 10.44.

Found: C, 44.72; H, 4.03; N, 10.32.

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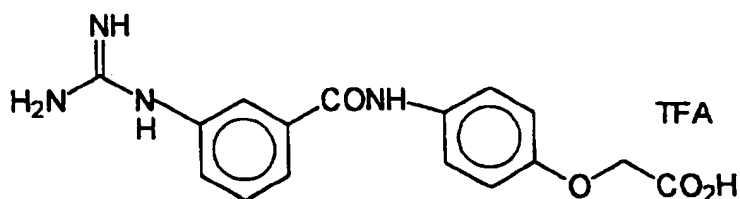
The compounds of the following Examples were prepared by hydrolyzing the compounds of Examples 89, 90 and 80 in the same manner as described in Example 35:

5

Example 81

Synthesis of [4-[[[3-[(aminoiminomethyl)amino]phenyl]-carbonyl]amino]phenoxy]acetic acid, trifluoroacetate salt

10



15

Analysis Calculated for C₁₆H₁₆N₄O₄ · 1 TFA · 0.9 H₂O:

C, 47.15; H, 4.13; N, 12.22.

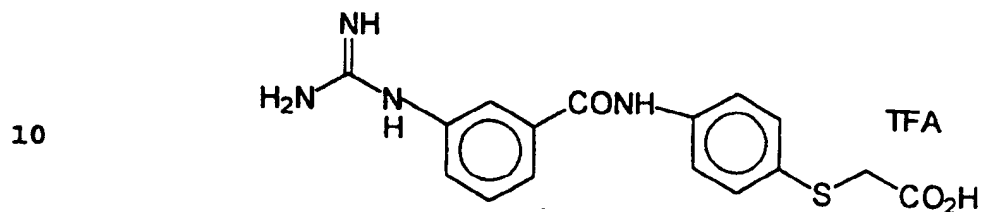
Found: C, 47.16; H, 3.85; N, 12.16.

20

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Example 82

Synthesis of [[4-[[[3-[(aminoiminomethyl)amino]phenyl]
carbonyl]amino]phenyl]thio]acetic acid,
5 trifluoroacetate



15 Analysis Calculated for C₁₆H₁₆N₄O₃S·1 TFA:
C, 47.16; H, 3.34; N, 12.22.
Found: C, 47.05; H, 3.61; N, 12.30.

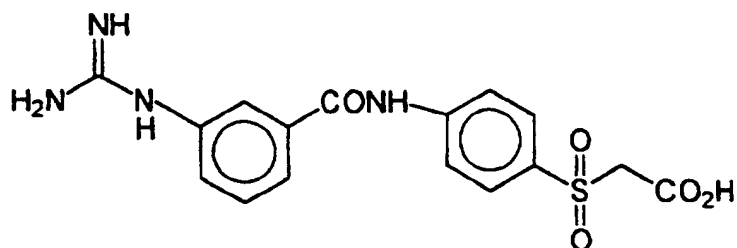
- 186 -

Example 83

Synthesis of [[4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]amino]phenyl]sulfonyl]acetic acid

5

10



Analysis Calculated for $C_{16}H_{16}N_4O_5S \cdot 0.6$ TFA:

15

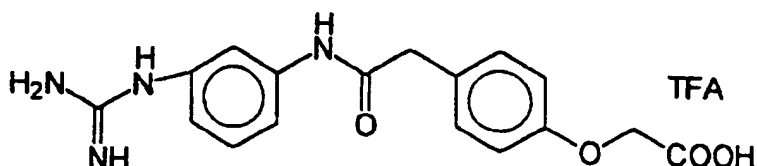
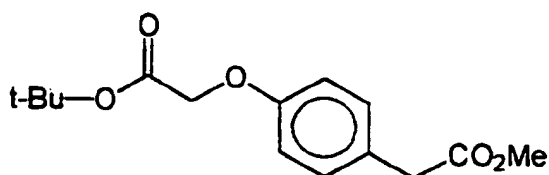
C, 46.44; H, 3.77; N, 12.60.

Found: C, 46.27; H, 3.69; N, 12.95

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Example 84

Synthesis of [4-[2-[[3-[(aminoiminomethyl)amino]phenyl]amino]-2-oxoethyl]phenoxy]acetic acid, trifluoroacetate salt

Step A

Methyl 4-hydroxyphenylacetate (2.0 g, 12.0 mmol, Aldrich) in THF (100 mL) was cooled (-30°C) and treated with sodium hydride (50% dispersion in mineral oil, 0.6 g, 12.2 mmol) in small portions over 15 minutes. The solution was then warmed (0°C) and stirred 30 minutes and then recooled to -30°C. To this solution was added neat t-butyl bromoacetate (2.6 g, 13.2 mmol, Aldrich) and the mixture was stirred 1 hour at -30°C and then warmed to room temperature and stirred 1 hour. The volatile components were removed at reduced pressure on a rotary evaporator and the residue was taken up in ether (50 mL). The ether was washed with water (25 mL), 10% NaOH (25 mL) and brine (25 mL). This produced 3.4 g (100%) of the title compound.

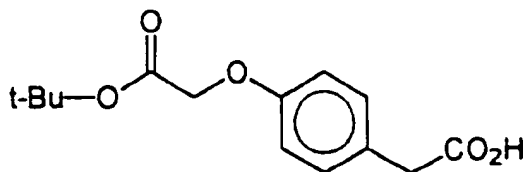
HRMS (M+) for C₁₅H₂₀O₅

Calculated: 280.1311

Found: 280.1297

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Step B



5

The product of Step A (0.5 g, 1.78 mmol) in THF (5 mL) was cooled (0°C) and treated with 1N lithium hydroxide solution (1.9 mL, 1.9 mmol). The solution was warmed to room temperature and stirred for 18 hours. The volatile components were removed at reduced pressure on a rotary evaporator. The crude product was chromatographed on silica gel gradient eluting with ethyl acetate:hexane (1:19 to 1:9 containing 1% acetic acid) and produced 0.16 g (25%) of the above compound.

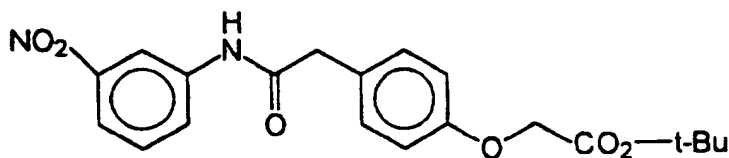
15

HRMS (M+) for $C_{15}H_{15}O_5$

Calculated: 266.1154

Found: 266.1163

Step C



20

The product from Step B (0.16 g, 0.6 mmol) and 3-nitroaniline (0.2 g, 1.4 mmol, Aldrich) in methylene chloride (5 mL) was cooled (0°C) and treated with 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (0.2 g, 1.04 mmol, Aldrich) and N-methylmorpholine (0.15 mL, 1.4 mmol, Aldrich). The solution was warmed to room temperature and stirred 18 hours. The mixture was poured into water and extracted with ethyl acetate (2 x 25 mL). The combined extracts were washed with water (2 x 10 mL) and saturated brine (10 mL), and dried over MgSO₄. The volatile components were removed at reduced pressure on a rotary evaporator to produce 0.19 g (82%) of the above compound.

35

HRMS (M+) for $C_{20}H_{22}N_2O_6$

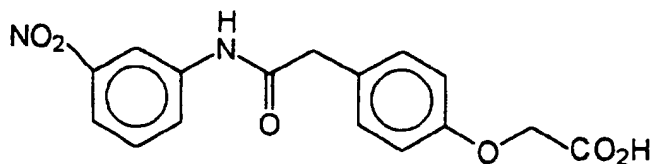
Calculated: 386.1478

Found: 386.1492

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Step D

5



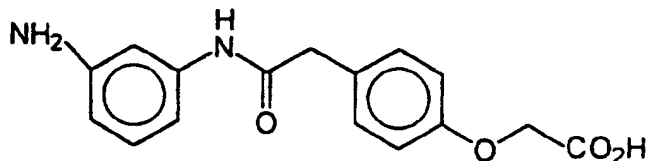
The product of Example 84C (0.19 g, 0.49 mmol) in methylene chloride (5 mL) was cooled (0°C) and treated with trifluoroacetic acid (0.5 mL). The solution was warmed to room temperature and stirred 3 hours. The volatile components were removed at reduced pressure on a rotary evaporator. This produced 0.14 g (86%) of the title compound.

15

HRMS (M+) for $C_{16}H_{14}N_2O_6$	Calculated:	330.0851
	Found:	330.0832

Step E

20



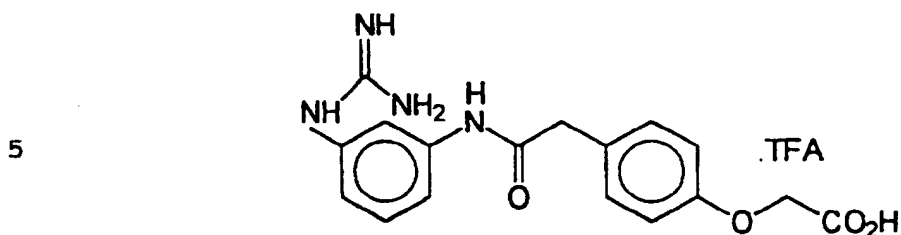
25

The product from Example 84D (0.14 g, 0.42 mmol) was subjected to the reaction conditions described for the preparation of Example 68B. This produced 0.11 g (88%) of the title compound.

30

HRMS (M+) for $C_{16}H_{16}N_2O_4$	Calculated:	300.1110
	Found:	

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Step F

A stirred solution of the product of Example 84E
10 (0.11 g, 0.37 mmol), diisopropylethylamine (0.09 mL)
and pyrazole-1-carboxamide hydrochloride (73 mg, 0.5
mmol) in dioxane (3 mL) and water (0.5 mL) was heated
at reflux for 3 hours. After cooling to room
temperature, the solvents were removed at reduced
15 pressure and the residue was chromatographed (reverse
phase HPLC, gradient elution with
water/acetonitrile/trifluoroacetic acid). This
produced 0.028 g (16%) of the title compound.

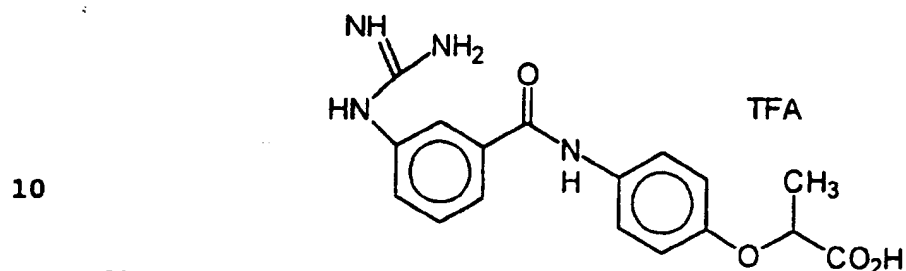
20 ESI MS (free base MH+) for $C_{17}H_{19}N_4O_4$

Calculated: 343

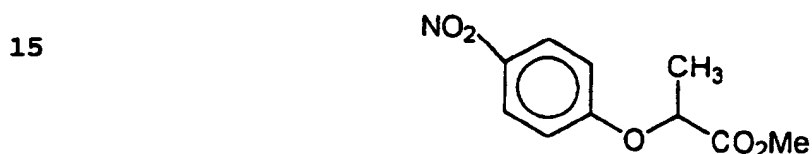
Found: 343

Example 85

Synthesis of 2-[4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]amino]phenoxy]propanoic acid, trifluoroacetate salt



Step A

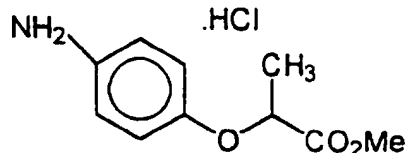


To a stirred and cooled (0°C) solution of 4-nitrophenol (0.56 g, 4 mmol, Aldrich), methyl DL-lactate (0.37 g, 3.6 mmol) and diethyl azodicarboxylate (0.64 mL, 4 mmol) in THF (50 mL) was added portionwise triphenylphosphine (1.05 g, 4 mmol). After 1 hour, the mixture was warmed to room temperature and stirred for 18 hours. The volatile components were removed at reduced pressure and the residue taken up in ethyl acetate (100 mL). The solution was washed with 5% aqueous sodium carbonate (4 x 25 mL), water (25 mL), brine (25 mL) and dried over magnesium sulfate. The volatile components were removed at reduced pressure on a rotary evaporator and the crude product was chromatographed on silica gel eluting with ether which produced 1.06 g of a 3:1 mixture of the above compound contaminated with 1,2-dicarbethoxyhydrazine.

HRMS (M+) for C₁₀H₁₁NO₅

Calculated:	225.0637
Found:	225.0636

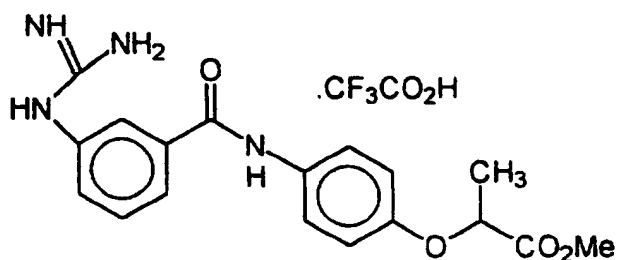
Step B



The crude product from Example 85A (1.06 g) was subjected to the reaction conditions described for the preparation of Example 68B. The crude product was
10 treated with methanolic HCl (5 mL) and then the volatile components were removed at reduced pressure. The residue was triturated with ether which produced 0.34 g of the title compound.

15	HRMS (free base M+) for C ₁₀ H ₁₃ NO ₃	Calculated: 195.0895
		Found: 195.0899

Step C



To a stirred and cooled (0°C) solution of 3-guanidinobenzoic acid hydrochloride (0.093 g, 0.43 mmol) and N-methylmorpholine (0.05 g, 0.43 mmol) in DMF (3 mL) was added isobutyl chloroformate (0.06 mL, 0.43 mmol). The mixture was stirred for 30 minutes. To this solution was added a solution of the product from Step B (0.1 g, 0.43 mmol) and N-methylmorpholine (0.05 g, 0.43 mmol) in DMF (2 mL). The reaction mixture was then allowed to warm to room temperature and stirred for 18 hours. The volatile components were removed at reduced pressure and the residue was chromatographed (reverse phase HPLC, gradient elution with

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water/acetonitrile/trifluoroacetic acid). This produced 120 mg (59%) of the above compound.

5 APCI MS (free base MH+) for $C_{18}H_{21}N_4O_4$ Calculated: 357
Found: 357

Step D

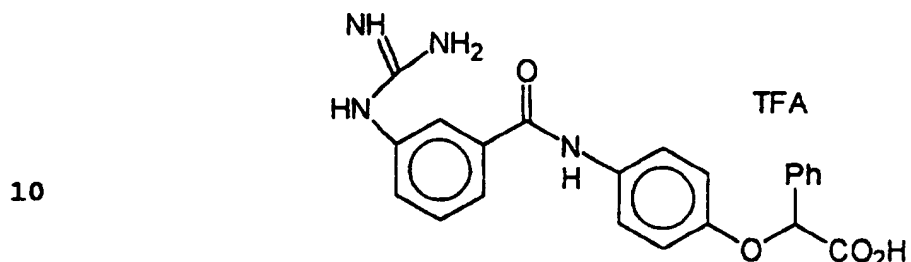
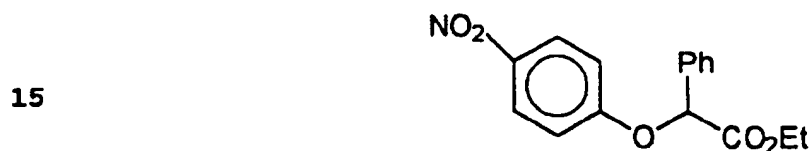
10 The product of Step C (0.05 g, 0.11 mmol) in methyl alcohol (1 mL) was cooled (0°C) and treated with 1N lithium hydroxide solution (0.22 mL, 0.22 mmol). The solution was warmed to room temperature and stirred for 18 hours. The volatile components were removed at reduced pressure on a rotary evaporator. The crude product was chromatographed (reverse phase HPLC,
15 gradient elution with water/acetonitrile/trifluoroacetic acid). This produced 50 mg (100%) of the above compound.

20 APCI MS (free base MH+) for $C_{17}H_{19}N_4O_4$ Calculated: 343
Found: 343

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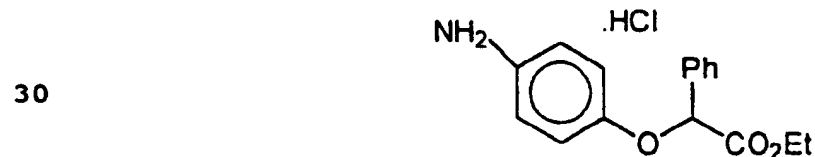
Example 86

Synthesis of α -[4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]amino]phenoxy]benzeneacetic acid,
5 trifluoroacetate salt

Step A

Ethyl DL-mandelate (0.65 g, 0.36 mmol) was
subjected to the reaction conditions described in
20 Example 85, Step A to produce 1.2 g of a mixture of the
above compound contaminated with 1,2-
dicarbethoxyhydrazine.

HRMS (M+) for $C_{16}H_{15}NO_5$ Calculated: 301.0950
25 Found: 301.0977

Step B

The crude product from Step A (1.2 g) in absolute
ethanol (10 mL) was treated with tin (II) chloride (2.4
35 g, 12.7 mmol) and then heated at 70°C for 1 hour. The
mixture was then cooled to room temperature and treated
with saturated sodium bicarbonate solution (25 mL) and

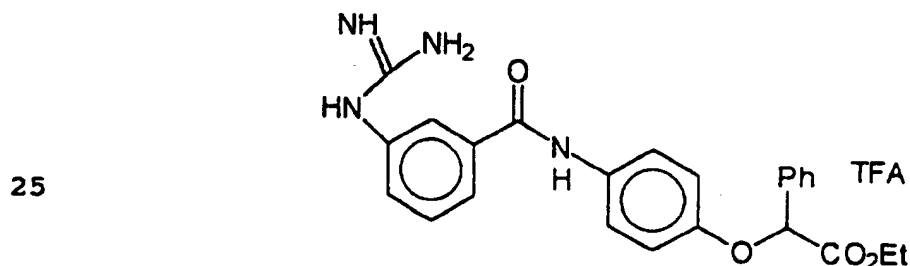
- 195 -

then poured into water (100 mL). The aqueous solution was then extracted with ethyl acetate (2 x 25 mL) and then poured into water (100 mL). The aqueous solution was then extracted with ethyl acetate (2 x 25 mL) and the combined extracts were washed with water (2 x 10 mL), brine (10 mL) and then dried over magnesium sulfate. The volatile components were removed at reduced pressure on a rotary evaporator. The residue was treated with methanolic HCl (5 mL) and the volatile components were removed at reduced pressure. The residue was triturated with ether which produced 0.55 g of the above compound.

15	HRMS (free base M+) for $C_{16}H_{17}NO_3$	Calculated: 271.1208
		Found: 271.1223

Step C

ethyl α -[4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]amino]phenoxy]benzeneacetate, trifluoroacetate
20 salt



The product from Step B (0.13 g, 0.43 mmol) was subjected to the reaction conditions described in Example 85, Step C to produce 0.11 g (47%) of the above compound.

APCI MS (free base MH+) for C₂₄H₂₅N₄O₄ Calculated: 433
Found: 433

35

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Step D

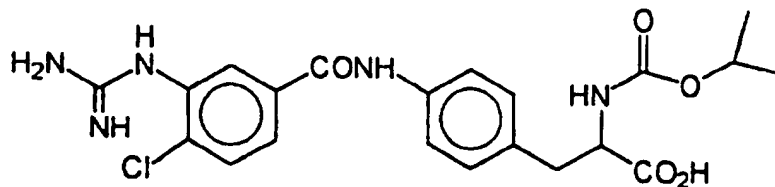
The product from Step C (0.05 g, 0.09 mmol) was subjected to the reaction conditions described in Example 85, Step D to produce 0.03 g (64%) of the above
5 compound.

APCI MS (free base MH+) for $C_{22}H_{21}N_4O_4$ Calculated: 405
Found: 405

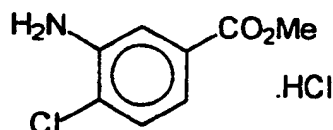
- 197 -

Example 87

4[[[3-[(aminoiminomethyl)amino]-4-chlorophenyl]-
carbonyl]amino]-N-[(1-methylethoxy)-
carbonyl]phenylalanine

Step AA

Preparation of

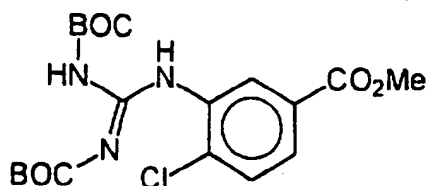


To a stirred suspension of 3-amino-4-chlorobenzoic acid (25.0 g, 157 mmol) in MeOH (300 ml) at 0°C, hydrogen chloride gas was added until the methanolic solution was saturated. The reaction mixture was stirred at 0-5°C for 30 minutes, allowed to attain room temperature, and then stirred for a further 4 days. The reaction mixture was concentrated *in vacuo* and the resulting white solid triturated with diethyl ether to afford the above compound; 26.2 g as a white solid.

NMR was consistent with the assigned structure.

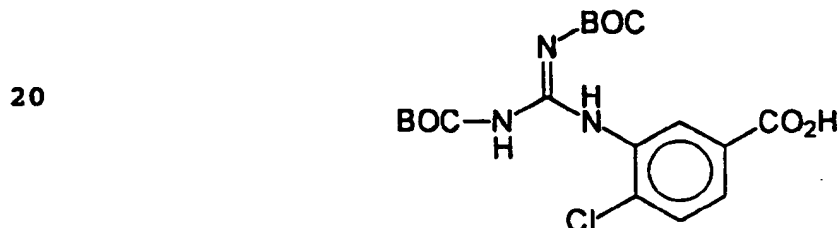
30 Step AAA

Preparation of



- 198 -

- To a solution of Example AZ and (24.8 g, 90 mmol) methyl-3-amino-4-chlorobenzoate (20 g, 90 mmol) in dimethylformamide (120 ml) and triethylamine (45 ml) at 0°C mercury II chloride (30.1 g, 111 mmol) was added.
- 5 The reaction mixture was stirred for 15 minutes at 0°C, allowed to attain room temperature and stirred for a further 2 hours. The reaction mixture was diluted with ethyl acetate (600 ml) and the resulting slurry filtered under reduced pressure. The filtrate was
- 10 concentrated, to afford an oily gum which was purified by chromatography on silica (eluent: ethyl acetate/heptane 20:80) to afford the above compound (8.6 g) as a white solid.
- 15 NMR was consistent with the assigned structure.

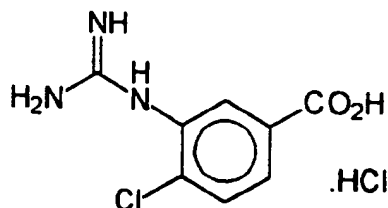
Step A

- 25 The product of Step AAA was dissolved in MeOH (3 mL) and 1 M NaOH (14 mL) was added at room temperature. The reaction was stirred at room temperature for 2 hours. The reaction was concentrated in vacuo and the residue dissolved in water, washed with ether. The
- 30 aqueous layer was acidified to pH=3 with 1N HCl. A white precipitate formed, was filtered and washed with water and ether and dried to give 1.2 g white solid. NMR was consistent with the proposed structure.

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Step B

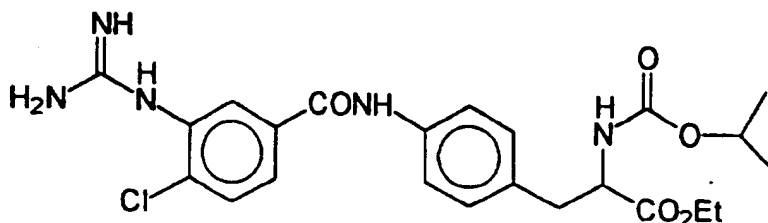
5



To a solution of the product of Step A (550 mg, 1.33 mmol) in CH_2Cl_2 (4 mL) was added TFA (1 mL) at 0°C .
10 The ice bath was removed after the addition and the reaction was stirred at room temperature for 2 hours. The reaction was concentrated in vacuo to give a colorless oil. To this was added 4N HCl solution in dioxane (2 mL) and white precipitate formed. The
15 solution was concentrated in vacuo to afford 280 g of the desired product as a white solid. NMR was consistent with the proposed structure.

Step C

20



25

A solution of the product of Step B (280 mg, 1.1 mmol) and 1-methyl piperidine (0.14 mL, 1.1 mmol) in DMF (5 mL) was cooled to 0°C and isobutyl chloroformate (0.14 mL, 1.1 mmol) was added under argon. The
30 reaction was allowed to stir for 5 minutes before adding a solution of Example AE (328 mg, 1.1 mmol) in DMF (2 mL). The reaction was allowed to warm slowly to room temperature over 16 hours. The solution was concentrated in vacuo and the residue purified by HPLC
35 to give the desired product as a yellow gummy oil (190 mg).

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Analysis Calculated for $C_{23}H_{28}N_5O_5Cl \cdot 1.5$ TFA:

C, 47.24; H, 4.50; N, 10.60.

Found: C, 47.23; H, 4.49; N, 10.97.

$M^+ = 489$.

5

Step D

The compound of Step C (170 mg, 0.34 mmol) was dissolved in MeOH (2 mL) and 1 M LiOH (1 mL) was added at room temperature. The reaction was stirred for 23 hours. The solution was concentrated in vacuo and the residue purified by HPLC to give the desired product as a white solid (130 mg).

10

Analysis Calculated for $C_{21}H_{24}N_5O_5Cl \cdot 1.4$ TFA:

15

C, 45.99; H, 4.12; N, 11.27.

Found: C, 45.95; H, 4.25; N, 11.29.

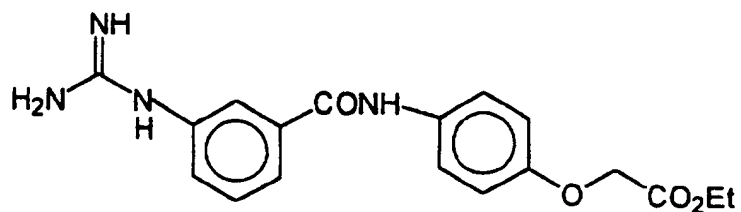
$M^+ = 461$.

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Example 89

Synthesis of ethyl [4-[[[3-[(aminoiminomethyl)amino]
phenyl]carbonyl]amino]phenoxy]acetate, trifluoroacetate
5 salt

10



15

The title compound was prepared under similar
conditions as described in Example 34, replacing the
compound of Example AA with the compound of Example BA.

Analysis Calculated for $C_{18}H_{20}N_4O_4 \cdot 1.1 \text{ TFA} \cdot 0.7 \text{ H}_2\text{O}$:

C, 49.07; H, 4.59; N, 11.33.

Found: C, 49.26; H, 4.23; N, 11.09.

20

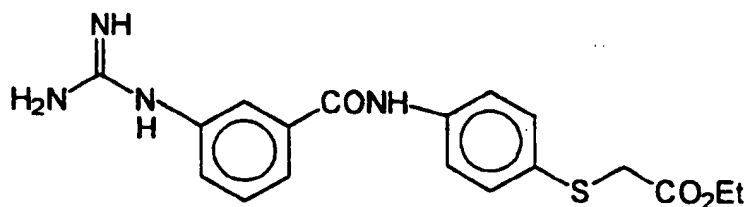
- 202 -

Example 90

Synthesis of ethyl[[4-[[[3-[(aminoiminomethyl)amino]-
phenyl]carbonyl]amino]phenyl]thio]acetic acid,
trifluoroacetate

5

10



15

The title compound was prepared in a similar
manner as described in Example 34, replacing the
compound of Example AA with the compound of Example BB.

Analysis Calculated for $C_{18}H_{20}N_4O_3S$:

C, 48.84; H, 4.43; N, 11.39.

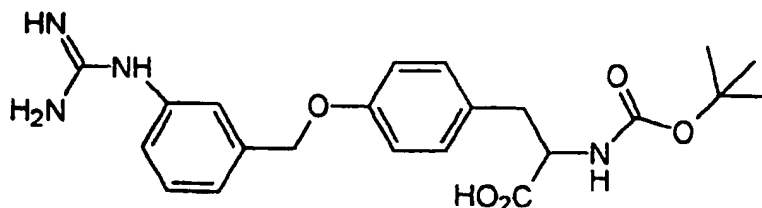
20

Found: C, 48.85; H, 4.12; N, 11.50.

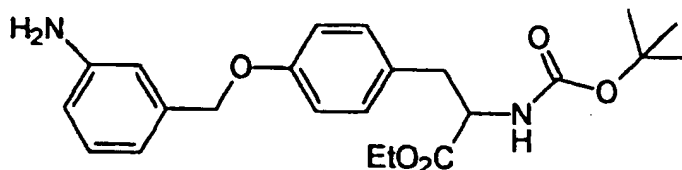
- 203 -

Example 91

Preparation of: 4-[[3-[(aminoiminomethyl)amino]-
phenyl]methoxy]-N-[(1,1-dimethylethoxy)-
carbonyl]phenylalanine



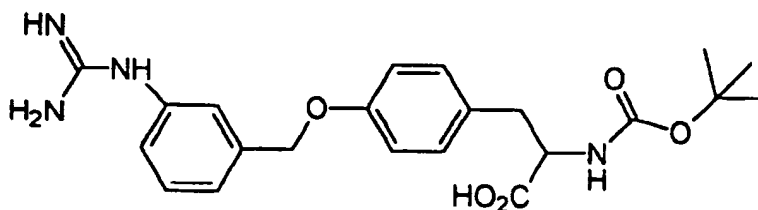
10 Step A



15

The N-t-BOC ethyl ester product of Example 11, Step A can be dissolved in absolute EtOH and reduced using substantially the quantities and procedure of Example 11, Step D to obtain the N-t-BOC aniline ethyl ester that can be isolated in substantially pure form by prep rphplc.

20 Step B



30

The N-t-BOC aniline ethyl ester of Step A can be converted to the guanidino - ethyl ester derivative using substantially the quantities, proportions and procedure of Example 11, Step E and subsequently hydrolyzed to the guanidino - acid derivative using the hydrolysis procedure of Example 11, Step E.

35

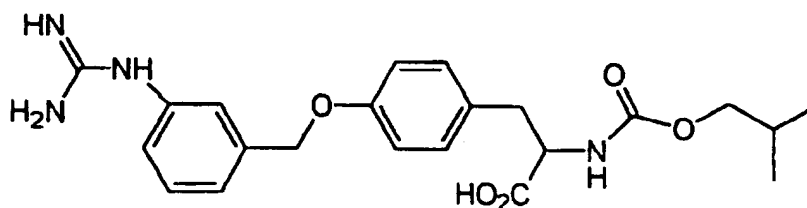
- 204 -

Example 92A

Preparation of: 4-[[3-[(aminoiminomethyl)amino]-
phenyl]methoxy]-N-[(1,1-dimethylethoxy)-
carbonyl]phenylalanine

5

10



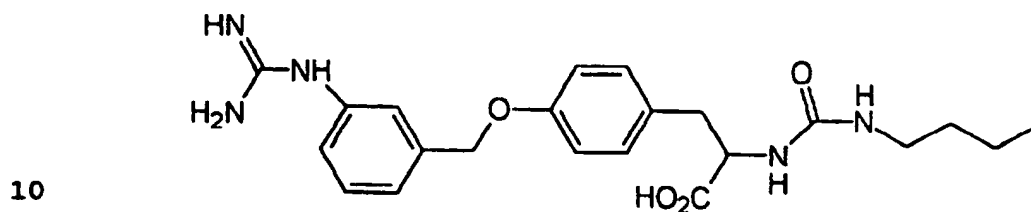
The title compound can be prepared using the
methodology of Example 11, substituting an equivalent
amount of isobutylchloroformate (Aldrich) for
n-butanesulfonyl chloride in Step C.

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Example 92B

Preparation of: 4-[[3-[(aminoiminomethyl)amino]phenyl]-
methoxy]-N-(butylamino)carbonyl]phenylalanine

5



The above compound can be prepared using the
methodology of Example 11, substituting an equivalent
amount of n-butyl isocyanate (Aldrich) for
15 n-butanesulfonyl chloride in Step C.

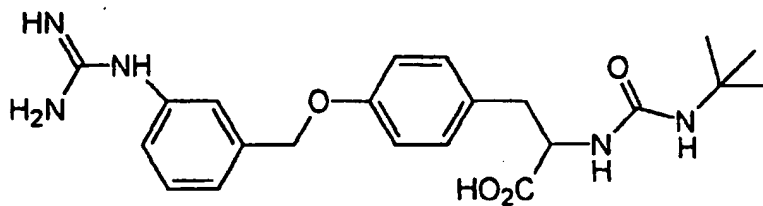
- 206 -

Example 93

Preparation of: 4-[[3-[(aminoiminomethyl)amino]-
phenyl]methoxy]-N-[[[(1,1-dimethylethyl)amino]-
carbonyl]phenylalanine

5

10



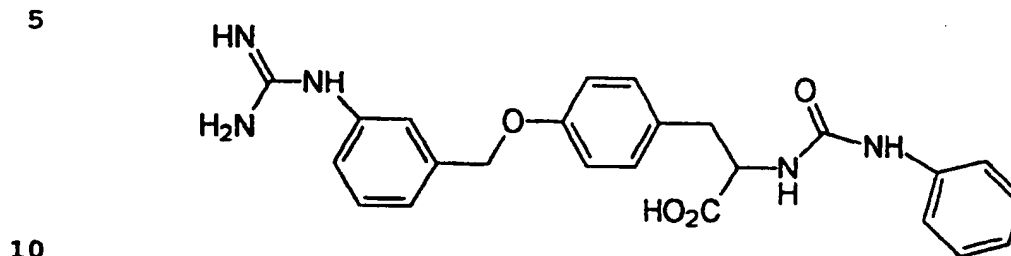
The above compound can be prepared using the
methodology of Example 11, substituting an equivalent
amount of t-butyl isocyanate (Aldrich) for
n-butanesulfonyl chloride in Step C.

15

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Example 94

Preparation of: 4-[[3-[(aminoiminomethyl)amino]phenyl]-methoxy]-N-(phenylamino)carbonyl]phenylalanine



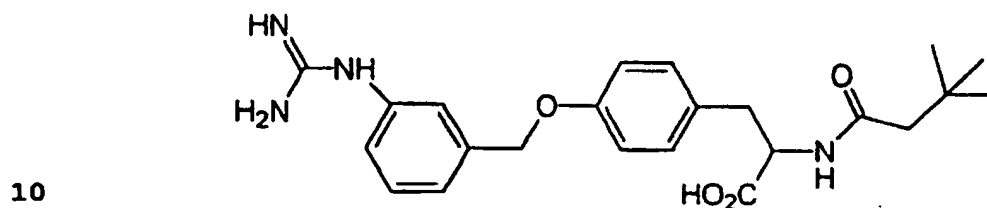
The above compound can be prepared using the methodology of Example 11, substituting an equivalent amount of phenyl isocyanate (Aldrich) for
15 n-butanesulfonyl chloride in Step C.

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Example 95

Preparation of: 4-[[3-[(aminoiminomethyl)amino]phenyl]-
methoxy]-N-(3,3-dimethyl-1-oxobutyl)phenylalanine

5



The above compound can be prepared using the
methodology of Example 11, substituting an equivalent
amount of isovaleryl chloride (Aldrich) for
15 n-butanesulfonyl chloride in Step C.

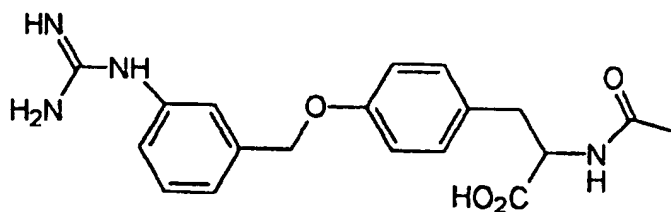
- 209 -

Example 96

Preparation of: N-acetyl-4-[[3-[(aminoiminomethyl)-
amino]phenyl]methoxy]phenylalanine

5

10



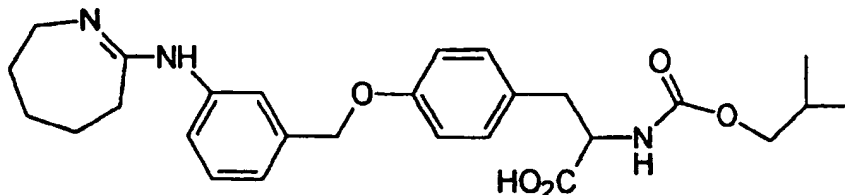
The above compound can be prepared using the methodology of Example 11, substituting an equivalent amount of acetyl chloride (Aldrich) for
n-butanesulfonyl chloride in Step C.

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Example 97

Preparation of: N-[(3-methylpropoxy)carbonyl]-4-[[3-
[(3,4,5,6-tetrahydro-2H-azepin-7-yl)amino]-
phenyl]methoxy]phenylalanine

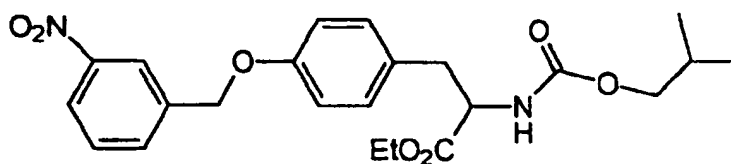
5



10

Step A.

15

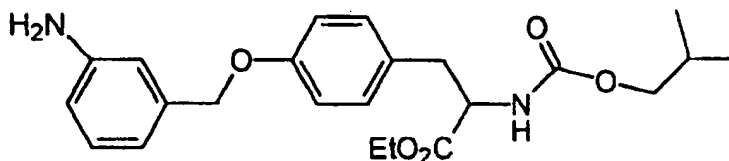


20

The above compound can be prepared using the procedure
of Example 11 and substituting an equivalent amount of
isobutyl chloroformate for butane sulfonylchloride in
Example 11, Step C.

Step B.

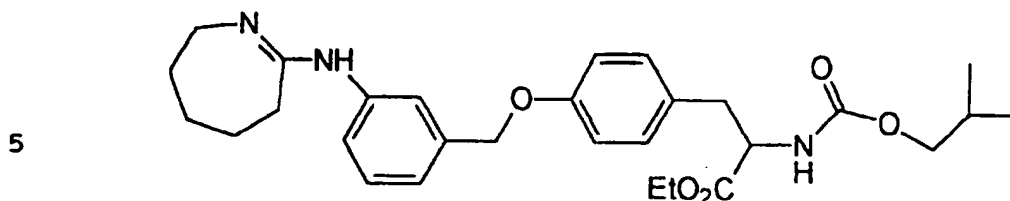
25



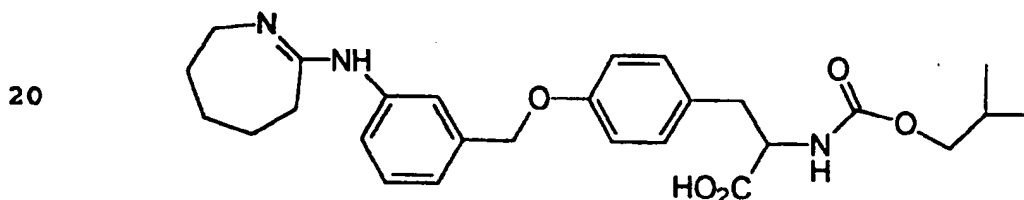
30

The nitro-tyrosine derivative of Step A can be
reduced to the aniline using substantially equivalent
amounts and the procedure of Example 11, Step D and
substituting the product of Step A for the product of
Example 11, Step C.

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Step C.

The above compound can be prepared by reacting essentially equivalent amounts of 1-aza-2-methoxy-1-cycloheptene with the anilino-tyrosine derivative prepared in Step B in a suitable solvent (EtOH, dimethylacetamide or dimethylformamide) until essentially complete reaction is obtained. The volatiles can be removed under reduced pressure and the desired product may be obtained by preparative RPHPLC.

Step D.

The ethyl ester obtained in Step C can be converted to the acid derivative using substantially the hydrolysis conditions of Example 11, Step E. The ethyl ester obtained in Step C can be dissolved in water:dioxane and aqueous lithium hydroxide added to pH 11. The reaction can be monitored by analytical RPHPLC until complete. The product can then be isolated by preparative RPHPLC.

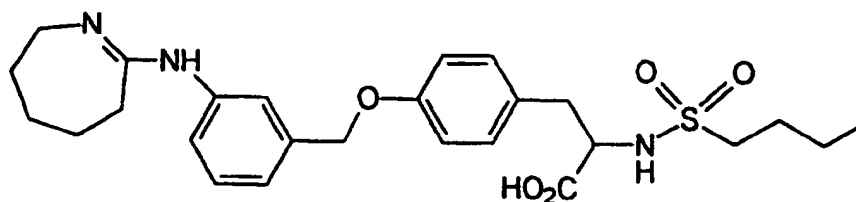
- 212 -

Example 98

Preparation of: N-(butylsulfonyl)-4-[[3-[(3,4,5,6-tetrahydro-2H-azepin-7-yl)amino]-phenyl]methoxy]phenylalanine

5

10



15

The above compound can be prepared using the procedures of Example 97 and using the procedure of Example 11, Step C to prepare the starting material (m-nitrobenzyl-N-n-butanesulfonyl tyrosine derivative). The product can be isolated using preparative RPHPLC.

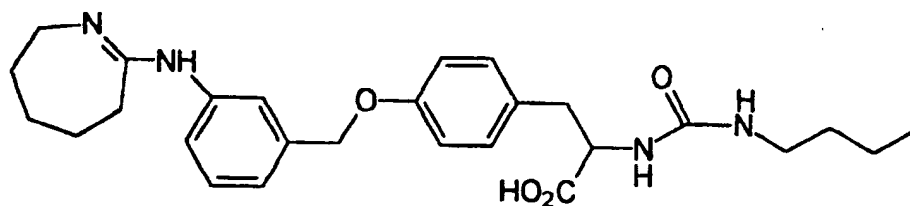
20

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Example 99

Preparation of: N-(butylamino)carbonyl]-4-[[3-
[(3,4,5,6-tetrahydro-2H-azepin-7-yl)amino]-
phenyl]methoxy]phenylalanine

5

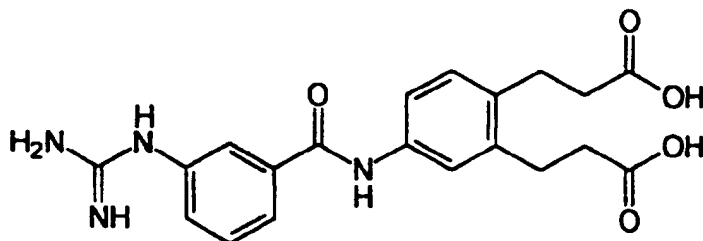
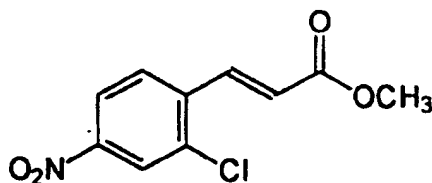


The above compound can be prepared using the
procedures of Example 97 and substituting n-butyl
isocyanate for n-butanesulfonyl chloride in Example 11,
10 Step C. The product can be isolated using preparative
RPHPLC.

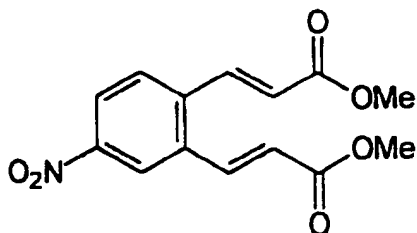
- 214 -

Example 100

4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]amino]benzene-1,2-dipropionic
acid, trifluoroacetate salt

Step A

To 2-chloro-5-nitro cinnamic acid (912 mg, 4 mmol)
in DMF (4 ml) was added NaHCO_3 (1.2 g) and MeI (1.1 g,
8 mmol) and the mixture stirred at room temperature
overnight. The solvent was removed and the residue
dissolved in CHCl_3 . The solution was washed with
water, dried and concentrated to provide the ester (800
mg). ^1H NMR was consistent with the proposed
structure.

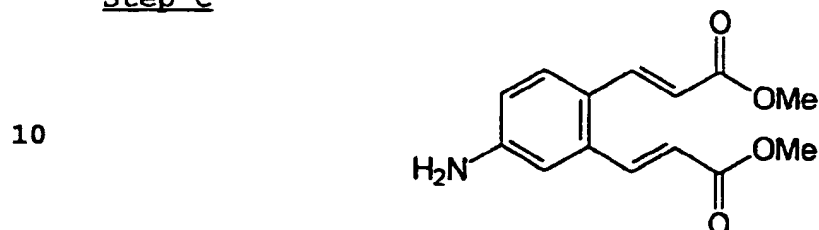
Step B

Methyl 2-chloro-5-nitrocinnamate acid (0.8 g, 3.3
mmol), methyl acrylate (450 mg, 5.2 mmol), tetra-*n*-
butylammonium chloride (920 mg, 3.3 mmol), NaHCO_3 (880

- 215 -

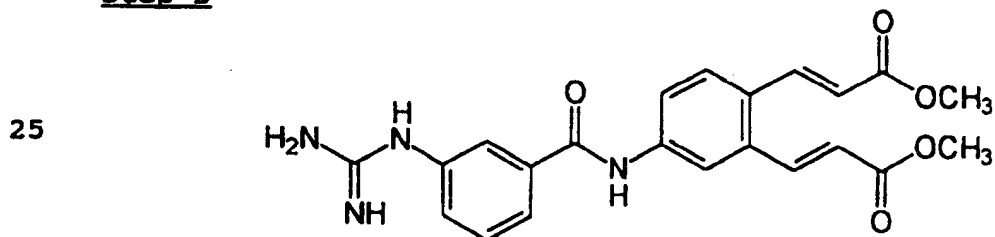
mg) and cat. $\text{Pd}(\text{OAc})_2$ (40 mg) were stirred in DMF (16 mL) at 80°C overnight. The mixture was cooled and the solvent removed. Flash chromatography (toluene/EtOAc) provided of the desired product (620 mg). ^1H NMR was
5 consistent with the proposed structure.

Step C



The product from Step B (620 mg, 2.1 mmol) and tin(II) chloride dihydrate (675 mg, 3 mmol) were stirred in 2:1 MeOH/concentrated HCl (19 ml) at 80°C for 1 hour. The mixture was cooled and filtered. The filtrate was concentrated and purified by flash chromatography (EtOAc/toluene/NH₄OH) to provide the desired compound (310 mg). ¹H NMR was consistent with the proposed structure.

Step D



To the compound of Example A (3-guanidinobenzoic acid hydrochloride, 300 mg, 1.4 mmol) in DMF (2 ml) at 0°C was added 1-hydroxybenzotriazole hydrate (HOBT, 200 mg, 1.5 mmol) and 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (EDCI, 300 mg, 1.6 mmol) and the mixture stirred at 0°C for 3 hours. The compound from Step C (308 mg, 1.2 mmol) was added and the mixture was stirred at room temperature for 48 hours. Purification by flash chromatography

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(MeOH/CHCl₃) and reverse phase HPLC (CH₃CN/H₂O/TFA) provided the desired compound. ¹H NMR and MS were consistent with the proposed structure.

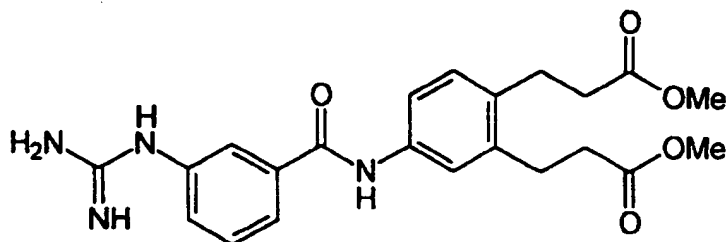
Analysis calculated for C₂₂H₂₂N₄O₅•1.5 TFA:

5 C, 50.60; H, 3.99; N, 9.44.

Found: C, 50.50; H, 4.18; N, 9.35.

Step E

10



15

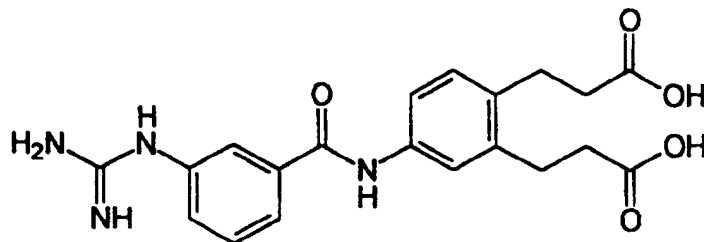
The product of Step D (300 mg) was hydrogenated at 5 psi H₂ in EtOH/MeOH/THF using 130 mg of 5% Pd/C as catalyst. Filtration, concentration, and purification by reverse phase HPLC (CH₃CN/H₂O/TFA) provided of the desired compound (260 mg). ¹H NMR and MS were

20

consistent with the proposed structure.

Step F

25



The product of Step E (200 mg, 0.47 mmol) was saponified as described in Example 35 to provide, after reverse phase HPLC (CH₃CN/H₂O/TFA), the desired compound. ¹H NMR and MS were consistent with the proposed structure.

30

Analysis calculated for C₂₀H₂₂N₄O₅•1.5 TFA:

35 C, 48.51; H, 4.16; N, 9.84.

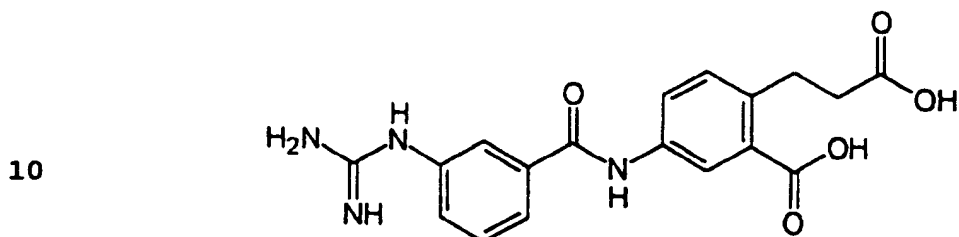
Found: C, 48.99; H, 4.20; N, 10.00.

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Example 101

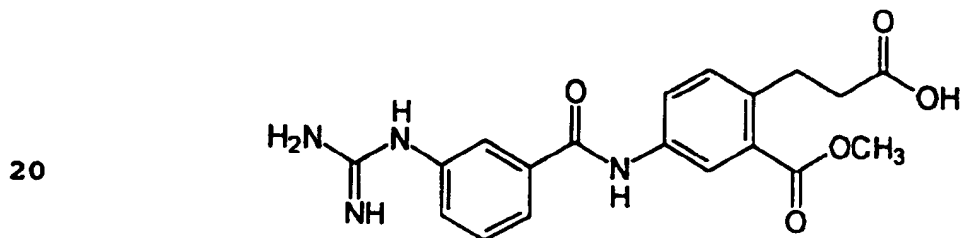
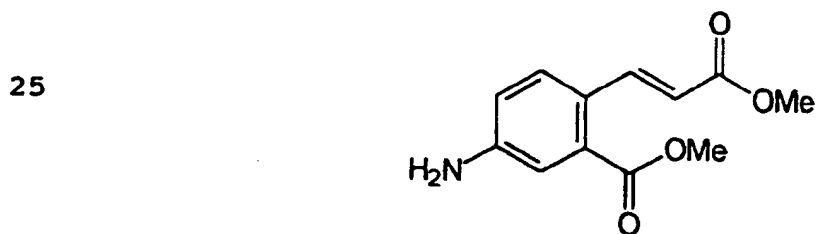
A:

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-2-carboxybenzenepropanoic acid,
5 bis(trifluoroacetate) salt



B:

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-2-methoxycarbonyl)benzenepropanoic acid,
15 bis(trifluoroacetate) salt

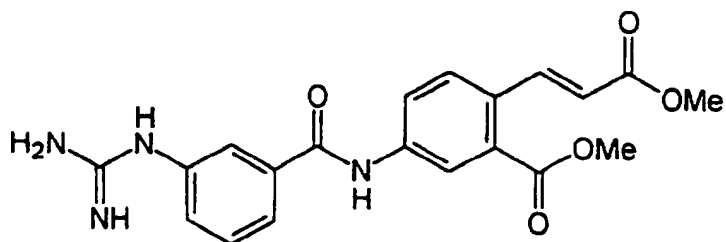
Step A/B

30 The desired compound was synthesized from methyl
2-bromo-5-nitrobenzoate using the procedure described
in Example 100, Steps B and C. ¹H NMR was consistent
with the proposed structure.

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Step C

5



10 The compound of Step A/B and 3-guanidinobenzoic acid hydrochloride were coupled under the conditions described in Example 76, Step D to provide, after reverse phase HPLC ($\text{CH}_3\text{CN}/\text{H}_2\text{O}/\text{TFA}$), the desired compound. ^1H NMR was consistent with the proposed structure.

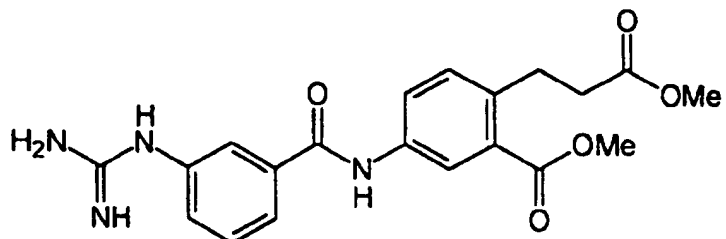
15 Analysis calculated for $\text{C}_{20}\text{H}_{19}\text{N}_4\text{O}_5 \cdot 1.5 \text{ TFA}$:

C, 48.77; H, 3.65; N, 9.89.

Found: C, 48.43; H, 3.88; N, 9.80.

Step D

20



25

The product of Step C (80 mg) was hydrogenated using the conditions described in Example 100 Step E. Purification by reverse phase HPLC ($\text{CH}_3\text{CN}/\text{H}_2\text{O}/\text{TFA}$) provided the desired compound (75 mg). ^1H NMR and MS were consistent with the proposed structure.

30

Analysis calculated for $\text{C}_{20}\text{H}_{24}\text{N}_4\text{O}_5 \cdot 2.0 \text{ TFA}$:

C, 48.77; H, 3.65; N, 9.89.

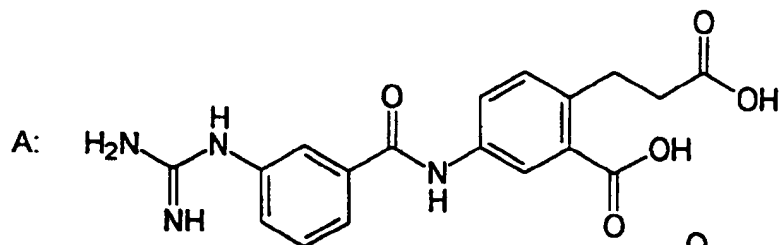
Found: C, 48.43; H, 3.88; N, 9.80.

35

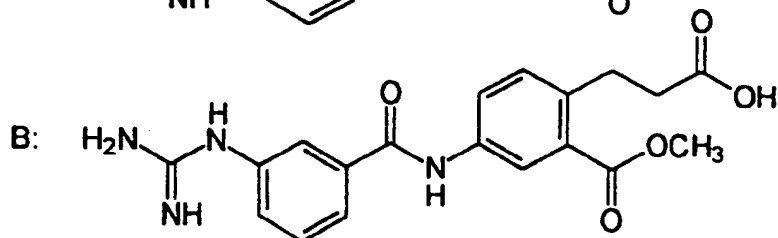
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Step E

5



10



The product of Step D (75 mg) was saponified as described in Example 35. Purification by reverse phase HPLC (CH₃CN/H₂O/TFA) provided 32 mg of A and 36 mg of B. ¹H NMR was consistent with the proposed structures

A: Analysis calculated for C₁₈H₁₈N₄O₅•2.2 TFA:

C, 43.31; H, 3.28; N, 9.02.

20

Found: C, 43.22; H, 3.56; N, 9.03.

B: Analysis calculated for C₁₉H₂₀N₄O₅•2.0 TFA:

C, 45.11; H, 3.62; N, 9.15.

Found: C, 45.38; H, 3.18; N, 9.43.

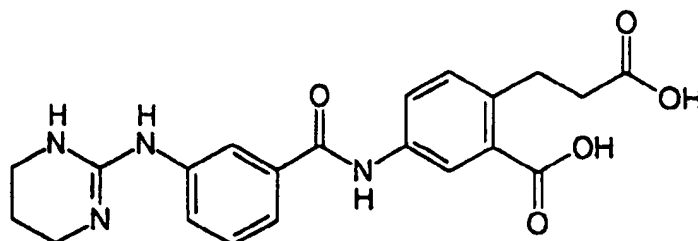
- 220 -

Example 102

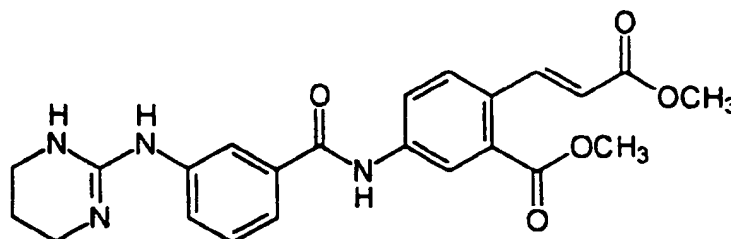
2-carboxy-4-[[[3-[(1,4,5,6-tetrahydropyrimidin-2-yl)-
amino]phenyl]carbonyl]amino]benzenepropanoic
acid, trifluoroacetate salt

5

10

Step A

15



20

The desired compound was prepared from the
compound of Example 101 Step A/B (270 mg, 1.1 mmol) and
the compound of Example 130 (280 mg, 1.1 mmol) using
the conditions described in Example 100, Step D.
Purification by flash chromatography (MeOH/CHCl₃)
provided the desired (270 mg). ¹H NMR and MS were
consistent with the proposed structure.

25

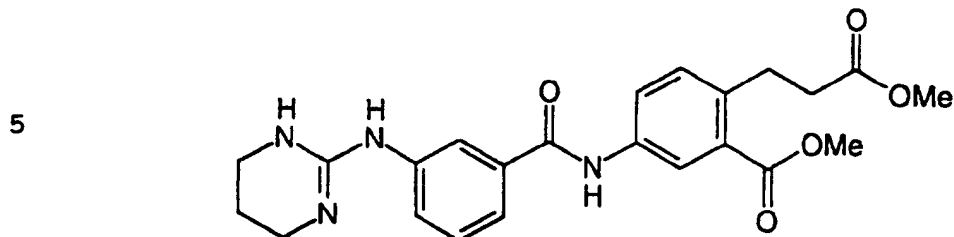
Analysis calculated for C₂₃H₂₄N₄O₅•1.2 TFA:

C, 53.22; H, 4.43; N, 9.77.

30

Found: C, 53.02; H, 4.20; N, 9.69.

- 221 -

Step B

10 The product of Step A (160 mg) was hydrogenated using the conditions described in Example 100 Step E, using MeOH, THF, TEA and NH_4OH to solubilize the starting material. Purification by flash chromatography ($\text{CHCl}_3/\text{MeOH}$) provided the desired
15 compound. ^1H NMR was consistent with the proposed structure.

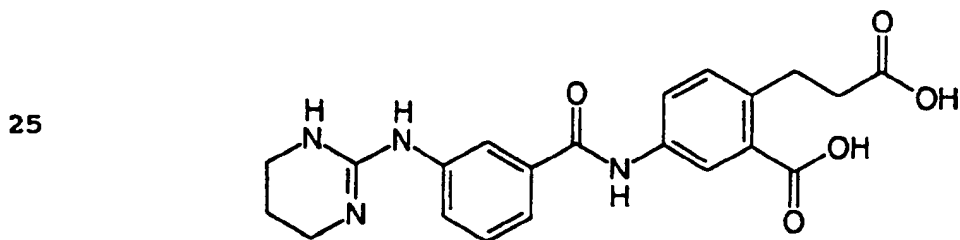
Analysis calculated for $\text{C}_{23}\text{H}_{26}\text{N}_4\text{O}_5 \cdot 1.0$ TFA:

C, 54.35; H, 4.93; N, 10.14.

Found:

C, 54.05; H, 5.21; N, 10.10.

20

Step C

30 The product of Step B (600 mg) was saponified as described in Example 35. Purification by reverse phase HPLC ($\text{CH}_3\text{CN}/\text{H}_2\text{O}/\text{TFA}$) provided the diacid (50 mg). ^1H NMR and MS were consistent with the proposed structure.

Analysis calculated for $\text{C}_{21}\text{H}_{22}\text{N}_4\text{O}_5 \cdot 1.3$ TFA:

C, 50.74; H, 4.20; N, 10.03.

35

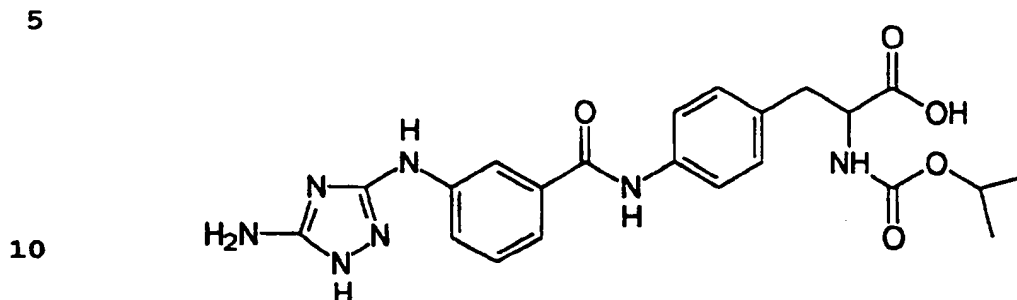
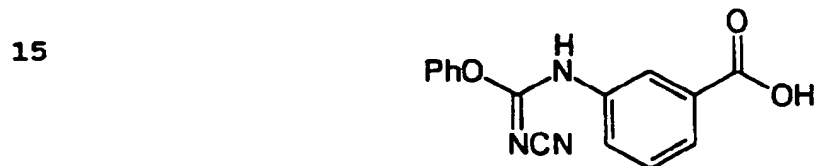
Found:

C, 50.72; H, 4.14; N, 9.93.

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Example 103

(±) 4-[[[3-[3-amino-1H-1,2,4-triazol-5-yl]amino]-phenyl]carbonyl]amino]-N-[(1-methylethoxy)carbonyl]phenylalanine, trifluoroacetate salt

Step A

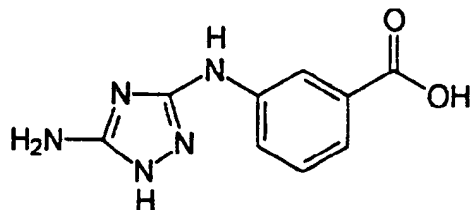
20 A solution of 3-aminobenzoic acid (2.0 g, 14.6 mmol) and diphenyl cyanocarbonimidate (3.48 g, 14.6 mmol) in isopropanol (25 mL) was stirred at room temperature for 18 hours. The resulting precipitate was filtered and washed with cold isopropanol and

25 recrystallized from acetonitrile to produce 1.7 g (41.4%) of the desired compound. ¹H NMR was consistent with the proposed structure.

HRMS (M ⁺) for C ₁₅ H ₁₁ N ₃ O ₃	calculated:	281.0800
	found:	281.0815

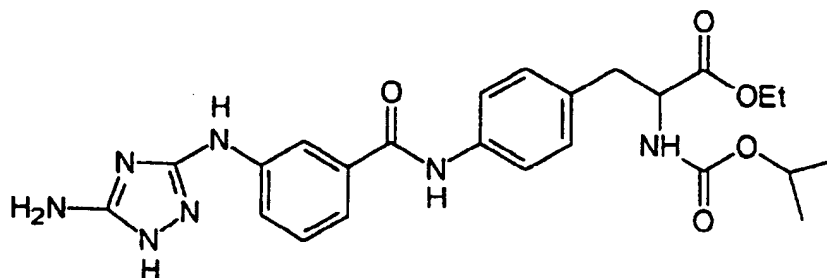
30

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Step B

The product from Step A (0.66 g, 2.35 mmol) and hydrazine hydrate (55% hydrazine) (0.15 g, 2.5 mmol) in methanol (20 mL) was stirred at room temperature for 18 hours. The resulting precipitate was filtered and washed with cold isopropanol to produce 0.6 g (77.4%) of the desired compound. ¹H NMR was consistent with the proposed structure.

HRMS (M+) for C ₉ H ₉ N ₅ O ₂	calculated:	219.0756
	found:	219.0749

Step C

The product from Step B (0.1 g, 0.6 mmol) and the product from Example AE (0.1 g, 0.45 mmol) in DMF (2.55 mL) were cooled to 0°C and treated with 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (0.09 g, 0.45 mmol) and N-methylmorpholine (0.05 mL, 0.45 mmol). The solution was warmed to room temperature and stirred 18 hours. The mixture was poured into water and extracted with ethyl acetate (2 x 25 mL). The combined extracts were washed with water (2 x 10 mL) and saturated brine (10 mL), and dried over MgSO₄. The volatile components were removed at reduced

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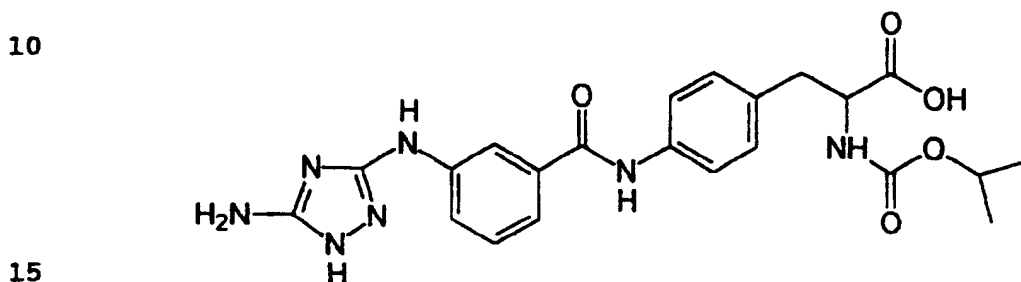
pressure on a rotary evaporator. The residue was purified by reverse phase HPLC (CH₃CN/H₂O/TFA) affording 70 mg (31.1 %) of the desired compound. ¹H NMR was consistent with the proposed structure.

5 HRMS (M+, free base) for $C_{24}H_{29}N_7O_5$

calculated: 495.2230

found: 495.2244

Step D



The product of Step C (0.05 g, 0.08 mmol) was saponified as described in Example 35 to provide, after reverse phase HPLC ($\text{CH}_3\text{CN}/\text{H}_2\text{O}/\text{TFA}$), 40 mg (87.5 %) of the title compound. ^1H NMR was consistent with the proposed structure.

APCI-MS (MH⁺, free base) for C₂₂H₂₆N₇O₅

calculated: 468

found: 468

25

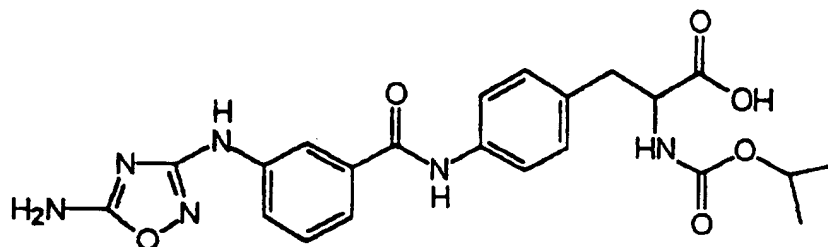
- 225 -

Example 104

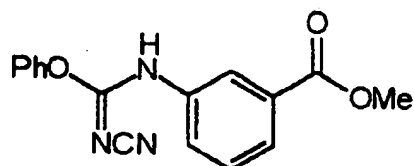
(±) 4-[[[3-[(5-amino-1,2,4-oxadiazol-3-yl)amino]-phenyl]carbonyl]amino]-N-[(1-methylethoxy)carbonyl]phenylalanine, trifluoroacetate salt

5

10

Step A

15



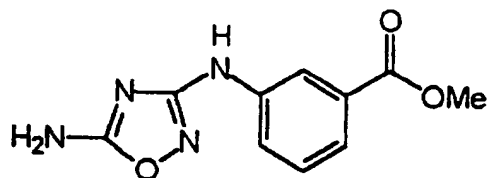
Methyl 3-aminobenzoate (2.2 g, 14.6 mmol, Lancaster) was subjected to the reaction conditions described in Example 103, Step A to produce 2.2 g (51%) of the desired compound. ¹H NMR was consistent with the proposed structure.

HRMS (M⁺) for C₁₅H₂₀O₅ calculated: 295.0957
found: 295.0957

25

Step B

30



The product from Step A (1.0 g, 3.39 mmol) in methanol (25 mL) was treated with a solution of hydroxylamine hydrochloride (0.24 g, 3.4 mmol), 50% sodium hydroxide (0.28 g, 3.4 mmol) and water (2 mL)

35

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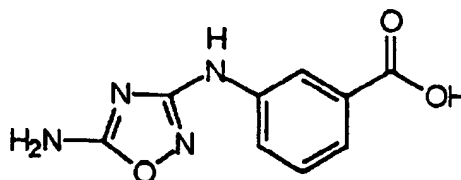
and the mixture stirred at room temperature for 18 hours. The mixture was concentrated and the resulting precipitate suspended in water, filtered and washed with cold water and cold isopropanol to produce 0.78 g (98%) of a mixture containing the desired compound along with an isomeric oxadiazole. ^1H NMR was consistent with the proposed structures.

HRMS (M+) for $\text{C}_{10}\text{H}_{10}\text{N}_4\text{O}_3$ calculated: 234.0753
found: 234.0752

10

Step C

15



20

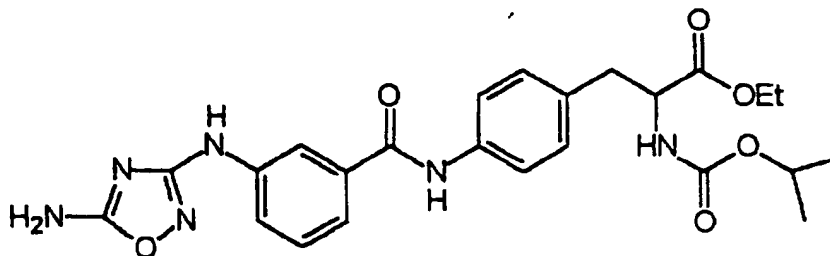
The product from Step B (0.71 g, 3.03 mmol) was saponified as described in Example 35, except that the mixture was heated at reflux for 4 hours, to produce 0.5 g (75%) of a mixture containing the desired compound along with an isomeric oxadiazole. ^1H NMR was consistent with the proposed structure.

25

HRMS (M+) for $\text{C}_{10}\text{H}_{10}\text{N}_4\text{O}_3$ calculated: 220.0596
found: 220.0605

Step D

30



35

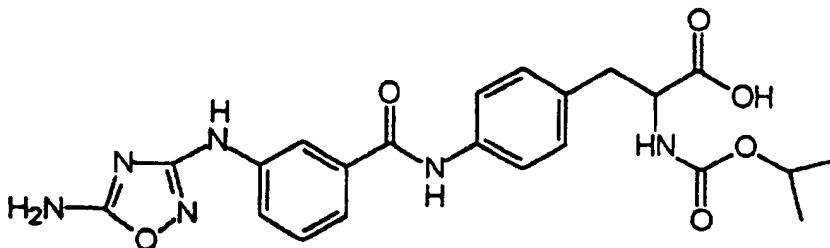
The product from Step C (0.1 g, 0.45 mmol) was coupled with the compound of Example AE as described in Example 103 Step C. The crude product was flash

- 227 -

chromatographed (methanol/chloroform/ammonium hydroxide) to produced 0.04 g (17.8%) of the desired compound (slower retention) and 0.02 g (8.9%) of the isomeric oxadiazole (faster retention). ¹H NMR was
5 consistent with the proposed structures.

HRMS (M+) for C ₁₀ H ₁₀ N ₄ O ₃	calculated:	496.2070
	found:	496.2069

Step E



The product from Step D (0.04 g, 0.08 mmol) was saponified as described in Example 35 to provide, after reverse phase HPLC ($\text{CH}_3\text{CN}/\text{H}_2\text{O}/\text{TFA}$), 30 mg (75%) of the title compound. ^1H NMR was consistent with the proposed structure.

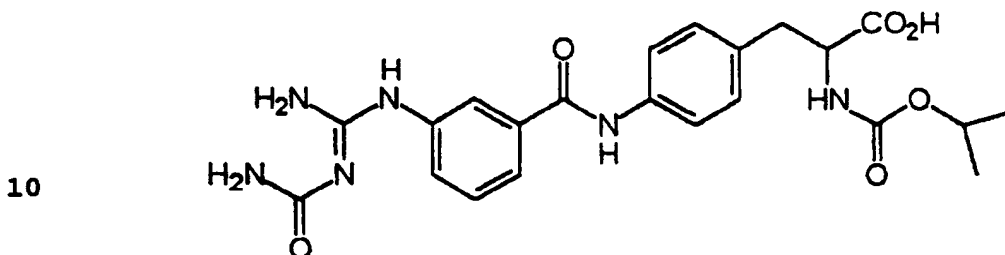
APCI-MS (MH+) for C₂₂H₂₅N₆O₆

calculated:	469
found:	469

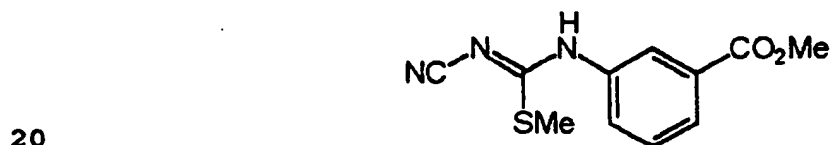
- 228 -

Example 105

4-[[[3-[[amino(aminocarbonyl)imino]methyl]-
amino]phenyl]carbonyl]amino]-N-[(1-
methylethoxy)carbonyl]phenylalanine,
bis(trifluoroacetate) salt

Step A

15 Preparation of methyl 3-[[[(cyanoimino)(methylthio)-
methyl]amino]benzoate



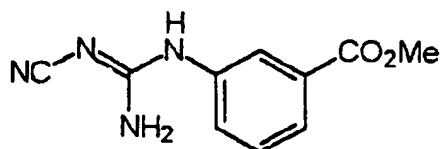
A stirred mixture of methyl 3-aminobenzoate (6.04 g, 40 mmol) and dimethyl N-cyanodithioiminocarbonate (11.96 g, 80 mmol) in pyridine (70 mL) was heated at
25 reflux for 2.5 hours. The reaction mixture was cooled to room temperature and upon standing overnight at room temperature the above compound (6.2 g) crystallized from the reaction mixture. The compound was used
30 without further purification. ¹H NMR was consistent with the proposed structure.

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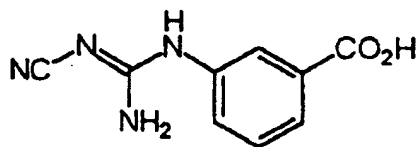
Step B

Preparation of methyl 3-[[amino(cyanoimino)methyl]-amino]benzoate

5



10 A mixture of the product from Step A (1.0 g) and ammonium hydroxide (2 mL) in ethanol (20 mL) was heated at 70°C in a sealed tube for 3.5 hours. The reaction mixture was cooled to room temperature and reduced to half its volume. After standing overnight at room
15 temperature a solid was obtained, which was filtered and washed with methanol to afford the above compound (389 mg) as a white solid. ¹H NMR was consistent with the proposed structure.

20 Step C

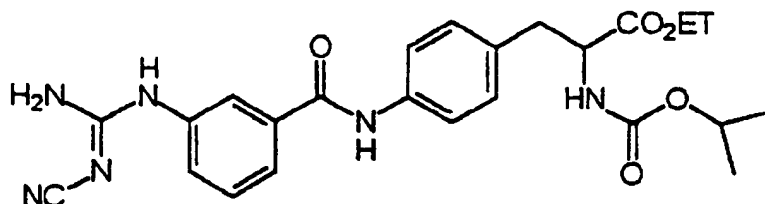
25

To a stirred solution of the product from Step B (2.9 g, 13.3 mmol) in THF (15 mL) and methanol (15 mL), was added 1N NaOH (14 mL). The reaction mixture was
30 stirred at room temperature for 2 hours and concentrated *in vacuo* to afford a white solid. The residue was acidified by suspension in water followed by addition of 1N HCl. The resultant solid was
35 filtered, washed with diethyl ether and dried to afford 2.4 g of the desired compound. ¹H NMR was consistent with the proposed structure.

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Step D

5

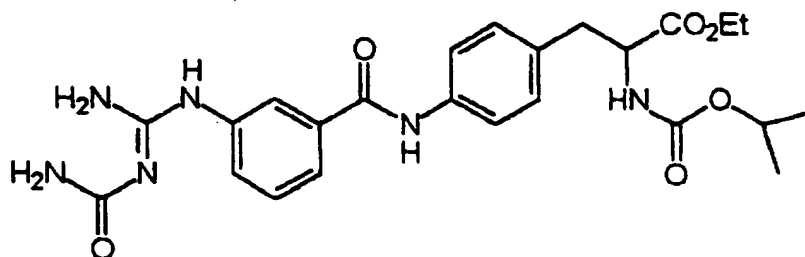


10 To a solution of the product from Step C (200 mg, 0.83 mmol) in DMF (1 mL) at -15°C was added N-methylmorpholine (0.10 mL, 0.83 mmol) and i-butylchloroformate (0.11 mL, 0.83 mmol). The product of Example AE (245 mg, 0.83 mmol) in DMF (0.5 mL) was
15 added and the reaction mixture stirred at -15°C for 30 minutes and at room temperature for 18 hours. The reaction mixture was concentrated *in vacuo* and the residue purified by flash chromatography (90:9:1 CH₂Cl₂/MeOH/NH₄OH) to give 398 mg of the desired
20 compound. ¹H NMR was consistent with the proposed structure.

Step E

25

30

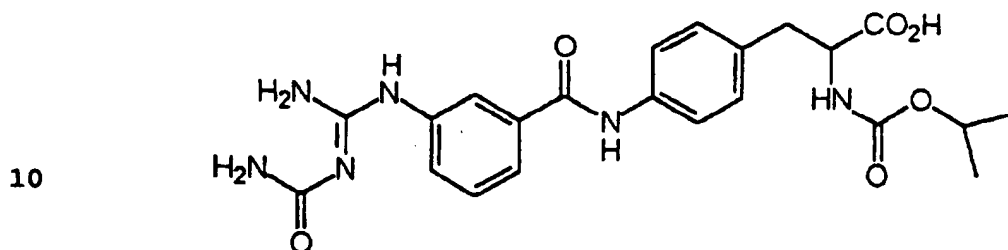


35 A solution of the product from Step D (95 mg, 0.20 mmol) in 1:1 CH₂Cl₂/TFA (1 mL) was stirred at room temperature for 2 hours. The solution was concentrated *in vacuo* and the residue purified by flash

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chromatography (90:9:1 CH₂Cl₂/MeOH/NH₄OH) to yield 65 mg of the desired compound. ¹H NMR was consistent with the proposed structure.

5 Step F

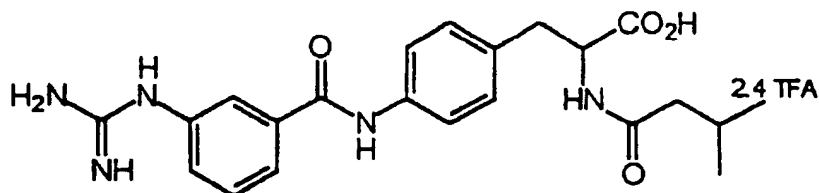
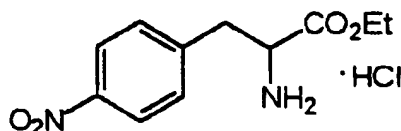


15 To a solution of the product from Step E (65 mg, 0.13 mmol) in 1:1 MeOH/THF (1 mL) was added 1N NaOH (0.5 mL). The mixture was stirred at room temperature for 2 hours and concentrated *in vacuo*. The residue was dissolved in water (1 mL) and the solution neutralized with 1N HCl. The white precipitate was collected and
20 purified by reverse phase HPLC (water/acetonitrile) to afford 15 mg of the above compound. ¹H NMR was consistent with the proposed structure.

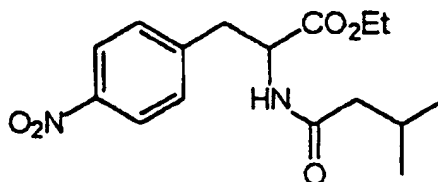
- 232 -

Example 106

4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]amino]-N-[(3-methyl-1-oxobutyl)-
phenylalanine, bis(trifluoroacetate) salt

Step A

HCl gas was bubbled through a solution of 4-nitro-DL-phenylalanine hydrate (5.00 g, 23.8 mmol) in absolute ethanol (80 ml) at 0°C for 5 minutes and the mixture refluxed overnight. The mixture was concentrated under reduced pressure and dried in vacuo to give a pale yellow solid (6.48 g, 98%). ¹H NMR was consistent with the proposed structure.

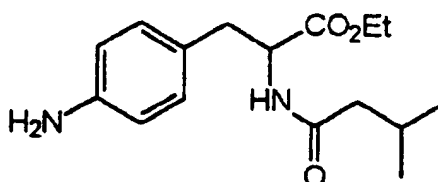
Step B

The product from Step A (2.00 g, 7.28 mmol) was dissolved in dichloromethane (50 ml) and cooled to 0°C. Isovaleryl chloride (1.23 g, 10.2 mmol) was added dropwise, followed by triethylamine (2.21 g, 21.8 mmol) and the mixture stirred overnight at room temperature. The mixture was concentrated and the residue dissolved

- 233 -

in ethyl acetate, washed with water and brine and dried over MgSO_4 . The solution was concentrated and dried in vacuo to give a pale yellow solid (2.30 g, 98%). ^1H NMR was consistent with the proposed structure.

5

Step C

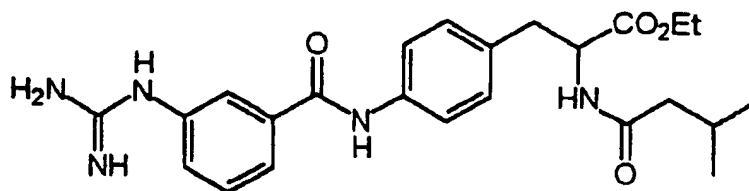
10

The product from Step B (2.30 g, 7.13 mmol) was dissolved in ethanol (30 mL) in a Parr vessel. 4% Pd/C (500 mg) was added and the mixture was hydrogenated at 5 psi for 4.5 hours. The solution was filtered through a plug of celite and the solution concentrated under reduced pressure. The residue was purified by flash chromatography on silica gel (60/40 ethyl acetate/heptane) to give a yellow solid (1.47 g, 70%). ^1H NMR was consistent with the proposed structure.

20

Step D

25



30

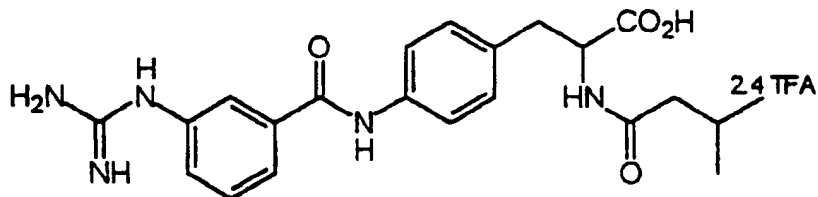
The product from Step C (439 mg, 1.5 mmol) and 3-guanidinobenzoic acid hydrochloride (323 mg, 1.5 mmol) were coupled under the conditions described in Example 76 Step D to provide, after purification by reverse-phase HPLC (water/acetonitrile), a white solid (190 mg, 28%). ^1H NMR was consistent with the proposed structure.

35

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Step E

5



The product from Step D (190 mg, 0.42 mmol) was
10 saponified as described in Example 35 to provide, after
purification by reverse-phase HPLC
(water/acetonitrile), the above compound as a white
solid (140 mg, 79%). ¹H NMR and MS were consistent
with the proposed structure.

15 Analysis calculated for C₂₂H₂₇N₅O₄•2.4 TFA:

C, 46.04; H, 4.24; N, 10.02.

Found: C, 46.03; H, 4.31, N, 9.85.

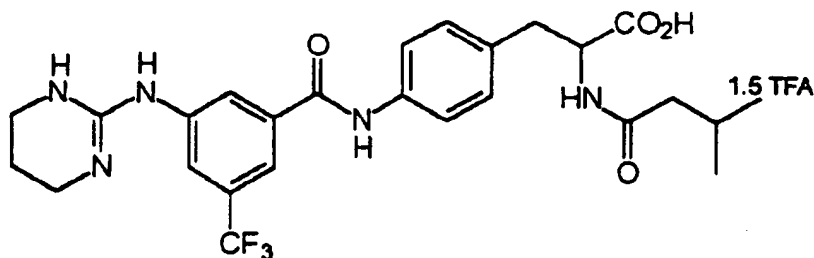
- 235 -

Example 107

N-(3-methyl-1-oxobutyl)-4-[[[3-[(1,4,5,6-tetrahydro-
pyrimidin-2-yl)amino]-5-(trifluoromethyl)phenyl]-
carbonyl]amino]phenylalanine, trifluoroacetate salt

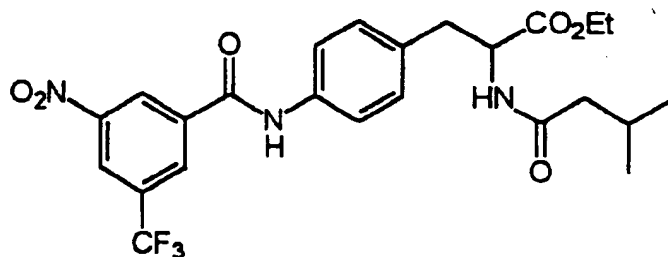
5

10

Step A

15

20



3-Nitro-5-trifluoromethylbenzoic acid (2.00 g, 8.51 mmol) was dissolved in toluene (20 ml) and cooled to 0°C. Oxalyl chloride (1.35 g, 10.63 mmol) was added to the solution, followed by 1 drop of DMF. The mixture was warmed to room temperature and stirred for 2 hours. The mixture was concentrated under reduced pressure to give the crude acid chloride as a clear yellow oil. A solution of the crude acid chloride (2.19 g, 8.64 mmol) in dichloromethane (20 ml) was added to a suspension of the product of Example 106 Step C (2.53 g, 8.64 mmol) in dichloromethane (20 ml). DMF (8 mL) was added to the suspension until homogeneous, followed by triethylamine (2.01 g, 19.9 mmol) and the mixture stirred at room temperature overnight. The mixture was concentrated and purified by flash chromatography on silica gel (70/30 ethyl

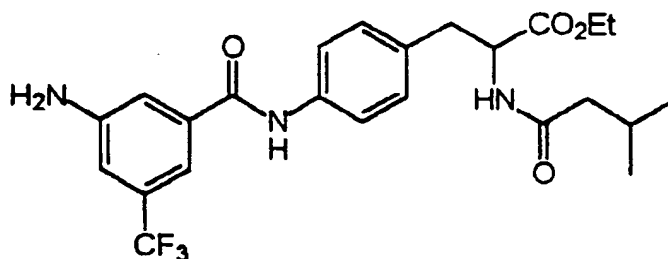
- 236 -

acetate/hexane) to give a yellow solid (3.79 g, 89%).
¹H NMR was consistent with the proposed structure.

Step B

5

10

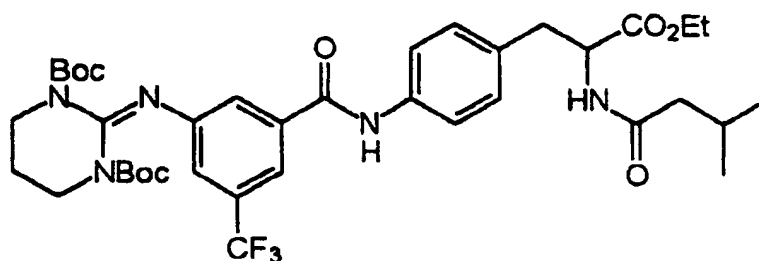


The product of Step A was reduced as described in
Example 106 Step C to provide the desired aniline. ¹H
NMR was consistent with the proposed structure.

Step C

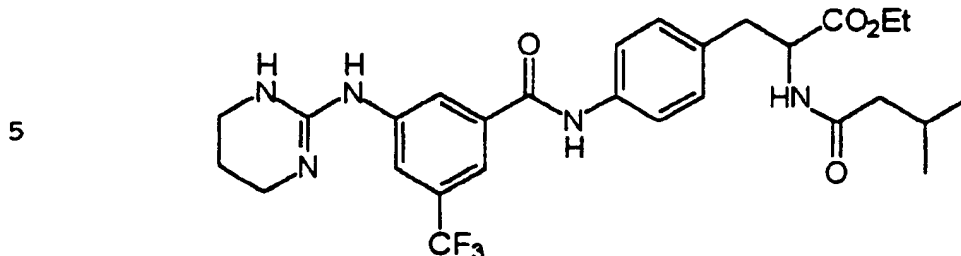
20

25



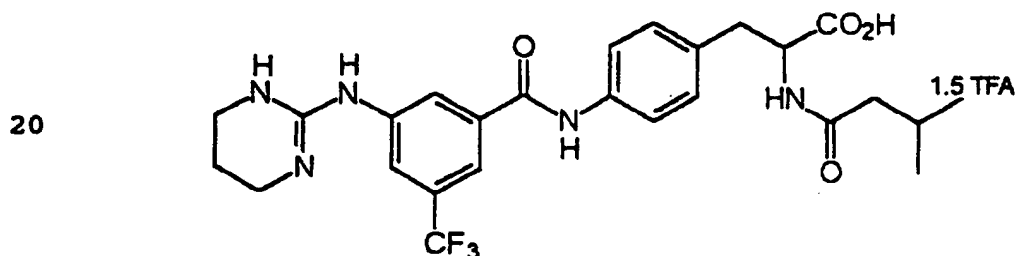
The above compound was synthesized from the
product of Step B (1.00 g, 2.09 mmol) and the product
of Example 131 (661 mg, 2.09 mmol) using the conditions
described in Example AS. Purification by flash
chromatography on silica gel (40/60 ethyl
acetate/hexane) gave a yellow solid (320 mg, 20%). ¹H
NMR was consistent with the proposed structure.

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Step D

10 The product of Step C (320 mg, 0.42 mmol) was treated with TFA as described in Example 60 to provide, after purification by reverse-phase HPLC (water/acetonitrile) a white solid (140 mg, 59%). ¹H NMR was consistent with the proposed structure.

15

Step E

25 The product of Step D (140 mg, 0.25 mmol) was saponified as described in Example 35 to provide, after purification by reverse-phase HPLC (water/acetonitrile) the above compound as a white solid (128 mg, 96%). ¹H NMR and MS were consistent with the proposed structure.

30 Analysis calculated for C₂₆H₃₀N₅O₄F₃•1.5 TFA:

C, 49.44; H, 4.51; N, 9.94.

Found: C, 49.64; H, 4.21; N, 9.87.

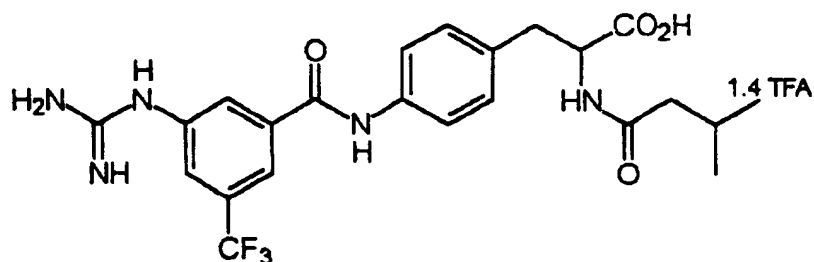
- 238 -

Example 108

4-[[[3-[(aminoiminomethyl)amino]-5-(trifluoromethyl)-
phenyl]carbonyl]amino]-N-(3-methyl-1-oxobutyl)
phenylalanine, trifluoroacetate salt

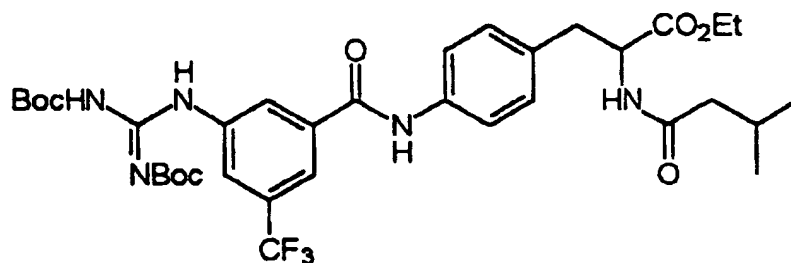
5

10

Step A

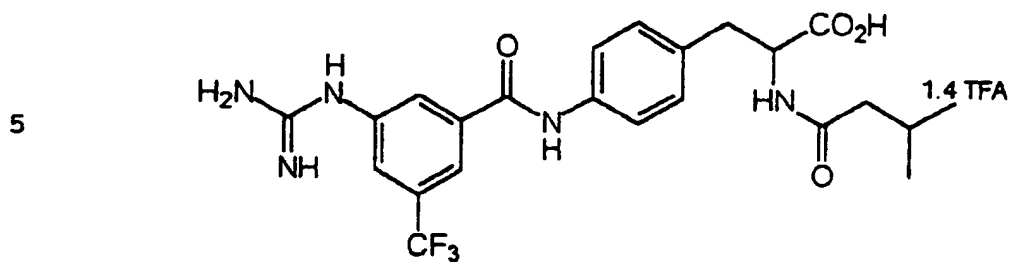
15

20



The compound from Example 107 Step B (2.016 g, 4.20 mmol) was treated with the product of Example AZ as described in Example AS to furnish, after
purification by flash chromatography (30/70 ethyl acetate/hexane) an off-white solid (960 mg, 32%). ¹H NMR was consistent with the proposed structure.

Steps B/C



10 The above compound was prepared from the product of Step A using the conditions described in Example 60 and Example 35. ¹H NMR and MS were consistent with the proposed structure.

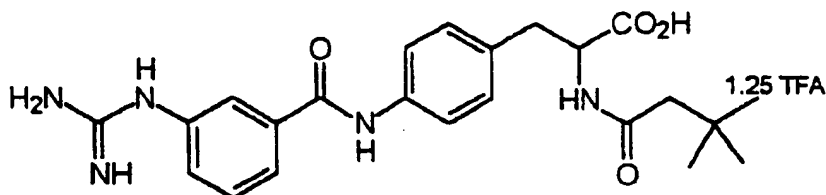
15 Analysis calculated for $C_{23}H_{26}N_5O_4F_3 \cdot 1.4$ TFA:
 C, 47.45; H, 4.23; N, 10.72.
 Found: C, 47.43; H, 3.96; N, 10.76.

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Example 109

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-(3,3-dimethyl-1-oxobutyl)phenylalanine,
trifluoroacetate salt

5



10

The above compound was prepared in the same manner
as described for the preparation of the compound of
Example 106, substituting t-butylacetyl chloride for
isovaleryl chloride. ¹H NMR and MS were consistent
with the proposed structure.

Analysis calculated for C₂₃H₂₉N₅O₄•1.25 TFA:

C, 52.62; H, 5.24; N, 12.03.

20

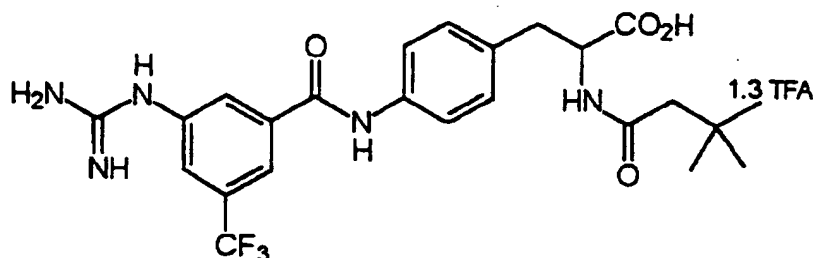
Found: C, 52.48; H, 5.49; N, 12.03.

- 241 -

Example 110

4-[[[3-[(aminoiminomethyl)amino]-5-(trifluoromethyl)-
phenyl]carbonyl]amino]-N-(3,3-dimethyl-1-oxobutyl)-
phenylalanine, trifluoroacetate salt

5



The above compound was prepared in the same manner
as described for the preparation of the compound of
15 Example 108, substituting t-butylacetyl chloride for
isovaleryl chloride. ¹H NMR and MS were consistent
with the proposed structure.

Analysis calculated for C₂₄H₂₈N₅O₄F₃•1.3 TFA:

C, 48.72; H, 4.50; N, 10.68.

20

Found:

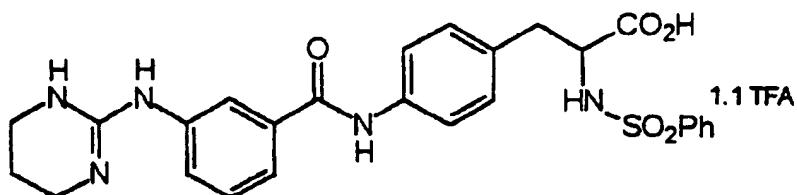
C, 48.54; H, 4.35; N, 10.72.

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Example 111

N-(phenylsulfonyl)-4-[[[3-[(1,4,5,6-tetrahydro-
pyrimidin-2-yl)amino]phenyl]carbonyl]amino]-
phenylalanine, trifluoroacetate salt

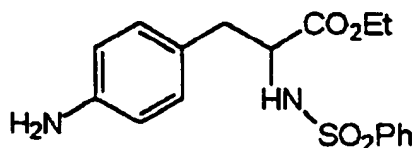
5



10

Steps A/B

15

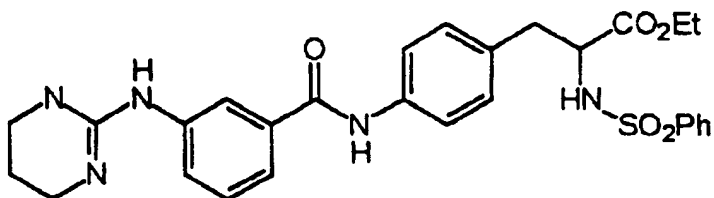


20

The above compound was prepared from the product
of Example 106 Step A using the procedures described in
Example 106 Steps B and C, substituting benzenesulfonyl
chloride for isovaleryl chloride. ¹H NMR was
consistent with the proposed structure.

Step C

25



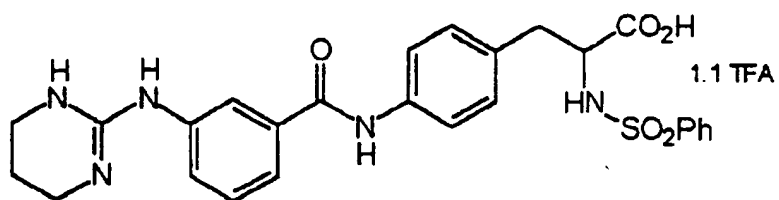
30

The desired compound was prepared from the product
of Step A/B and the product of Example 130, using the
procedure described in Example 106 Step D. ¹H NMR was
consistent with the proposed structure.

- 243 -

Step D

5



10 The product of Step C was saponified using the procedure described in Example 35 to provide the above compound. ¹H NMR and MS were consistent with the proposed structure.

Analysis calculated for C₂₆H₂₇N₅O₅S•1.1 TFA:

C, 52.35; H, 4.38; N, 10.82.

15

Found:

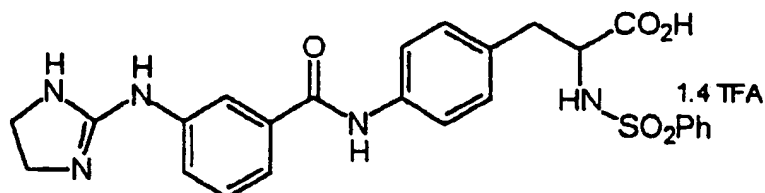
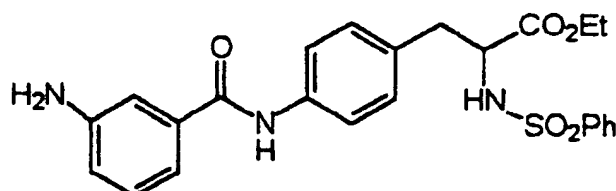
C, 52.42; H, 4.08; N, 10.91.

- 244 -

Example 112

4-[[[3-[(4,5-dihydro-1H-imidazol-2-yl)amino]phenyl]-
carbonyl]amino-N-(phenylsulfonyl)phenylalanine,
trifluoroacetate salt

5

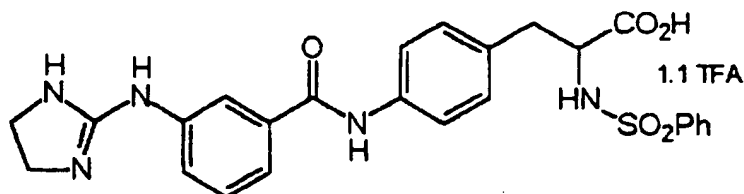
10 Steps A/B

15

The desired compound was prepared from the product
of Example 111 Step A/B and 3-nitrobenzoyl chloride
using the procedure described in Example 107 Steps A
and B. ¹H NMR was consistent with the proposed
20 structure.

Steps C/D/E

25



The above compound was prepared from the product
30 of Step A/B and the product of Example 132 using the
procedures described in Example 107 Steps C, D and E.
¹H NMR and MS were consistent with the proposed
structure.

Analysis calculated for C₂₅H₂₅N₅O₅S•1.4 TFA:

35

C, 50.05; H, 3.99; N, 10.50.

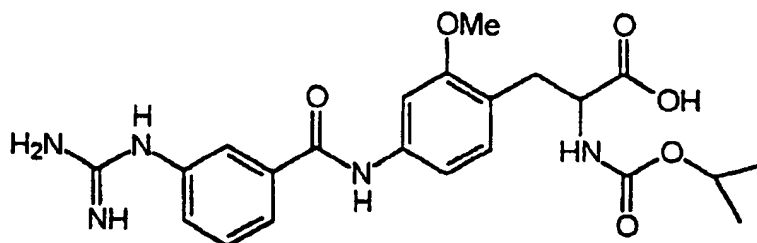
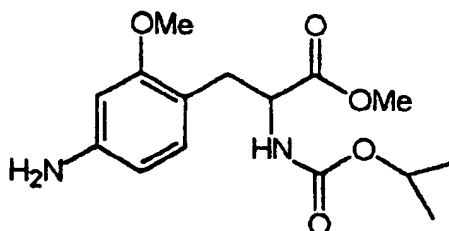
Found:

C, 49.88; H, 3.89; N, 10.62.

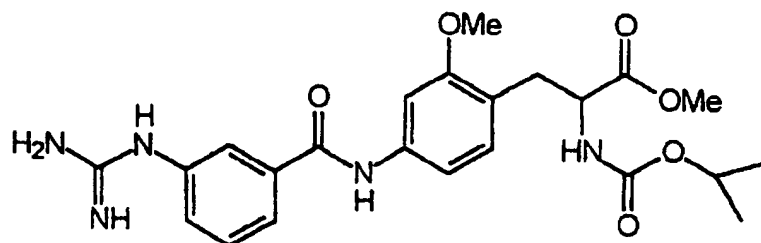
- 245 -

Example 113

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]amino]-
2-methoxy-N-[(1-methylethoxy)carbonyl]phenylalanine,
trifluoroacetate salt, monohydrate

Steps A/B/C

The above compound was prepared from methyl
pyruvate, isopropylcarbamate and 2-bromo-5-nitro
anisole using the procedures described in Example 76
Steps A, B and C.

Step D

The compound of Step A/B/C (370 mg, 1.19 mmol),
was coupled with 3-guanidinobenzoic acid hydrochloride
under the conditions described in Example 76 Step D to
give 450 mg of desired product. ¹H NMR was consistent
with the proposed structure.

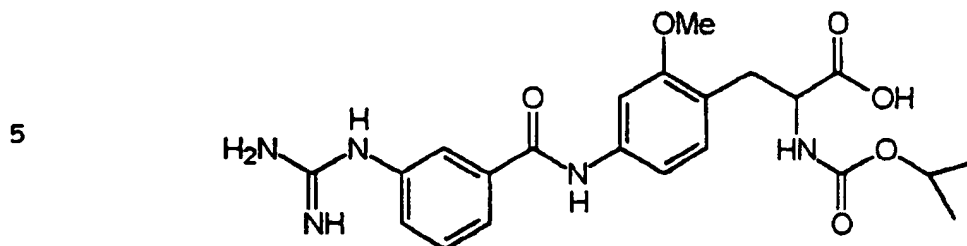
Analysis calculated for C₂₃H₂₉N₅O₆•1.0 TFA•0.8 H₂O:

C, 50.05; H, 5.31; N, 11.67.

Found:

C, 50.00; H, 4.91; N, 11.49.

Step E



10 The product of Step D (350 mg, 0.6 mmol) was saponified as described in Example 35 to give 230 mg of the above compound. ¹H NMR was consistent with the proposed structure.

Analysis calculated for $C_{22}H_{27}N_5O_6 \cdot 1.0 \text{ TFA} \cdot 1.0 \text{ H}_2O$:

15 C, 48.90; H, 5.13; N, 11.88.

Found: C, 48.67; H, 4.73; N, 11.76.

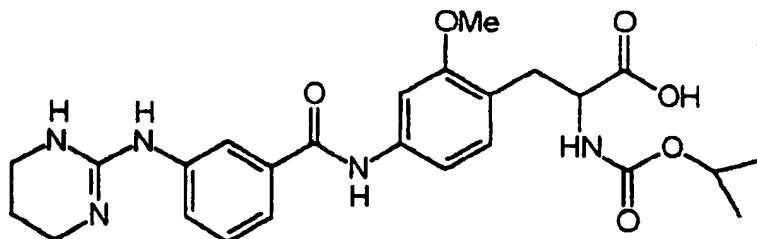
- 247 -

Example 114

2-methoxy-N-[(1-methylethoxy) carbonyl]-4-[[[3-
[(1,4,5,6-tetrahydropyrimidin-2-yl)amino]-
phenyl]-carbonyl]amino]phenylalanine,
trifluoroacetate salt

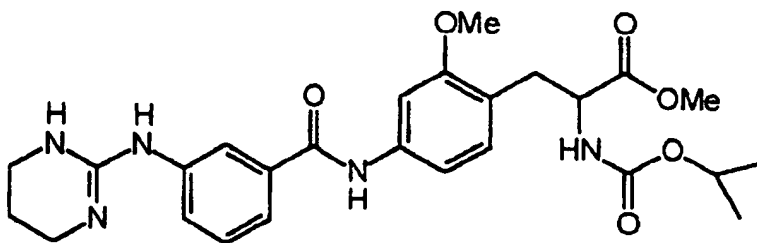
5

10

Step A

15

20



The above compound was prepared from the compound
of Example 113 Step A/B/C (485 mg, 1.56 mmol) and the
product of Example 130 (400 mg, 1.56 mmol) using the
conditions described in Example 76 Step D to give 710
mg of desired product. ^1H NMR was consistent with the
proposed structure.

Analysis calculated for $\text{C}_{26}\text{H}_{33}\text{N}_5\text{O}_6 \cdot 1.0 \text{ TFA}$:

C, 53.76; H, 5.48; N, 11.19.

30

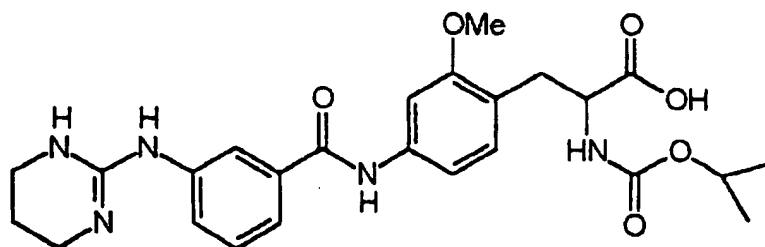
Found:

C, 53.54; H, 5.65; N, 11.11.

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Step B

5



10 The product of Step A was saponified as described
in Example 35 to give the above compound. ¹H NMR was
consistent with the proposed structure.

Analysis calculated for C₂₅H₃₁N₅O₆•1.0 TFA•0.5 H₂O:

C, 52.26; H, 5.36; N, 11.29.

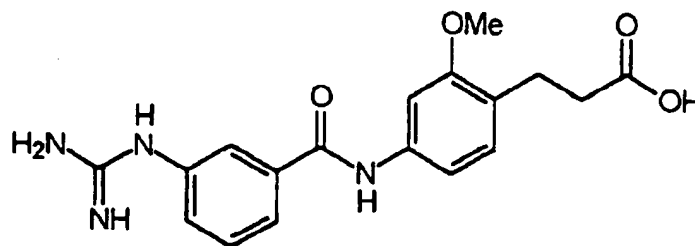
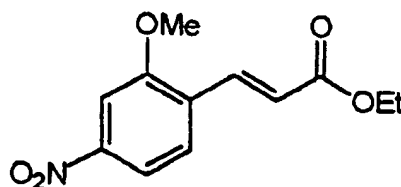
15

Found: C, 51.97; H, 5.08; N, 11.16.

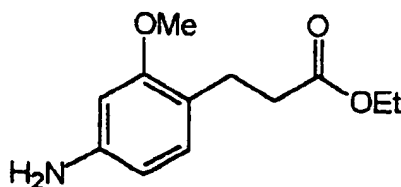
- 249 -

Example 115

4-[[[3-(aminoiminomethyl)amino]phenyl]-
carbonyl]amino]-2-methoxybenzenepropanoic
acid, trifluoroacetate salt

Step A

2-bromo-5-nitroanisole (1 g, 4.3 mmol) and ethyl
acrylate were coupled using the procedure described in
Example 100 Step B to give 1.10 g of the desired
compound. ^1H NMR was consistent with the proposed
structure.

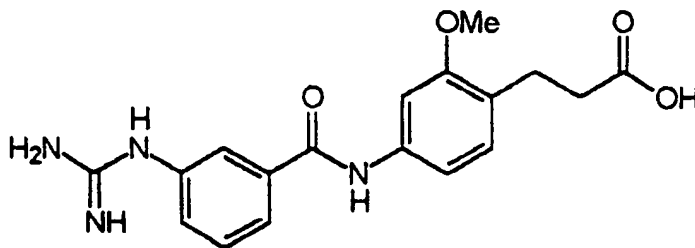
Step B

The compound of Step A (2 g, 0.85 mmol) was
reduced under the conditions described in Example 106
Step C to give, after flash chromatography (30/70 ethyl
acetate/hexane), 1.6 g of desired product. ^1H NMR was
consistent with the proposed structure.

- 250 -

Steps C/D

5



10

The compound from Step B was coupled with 3-guanidinobenzoic acid hydrochloride under the conditions described in Example 76 Step D and saponified as described in Example 35 to give the above compound. ¹H NMR was consistent with the proposed structure.

15

Analysis calculated for C₁₈H₂₀N₄O₄•1.4 TFA•1.4 H₂O:

C, 46.16; H, 4.51; N, 10.35.

Found:

C, 46.38; H, 4.36; N, 10.20.

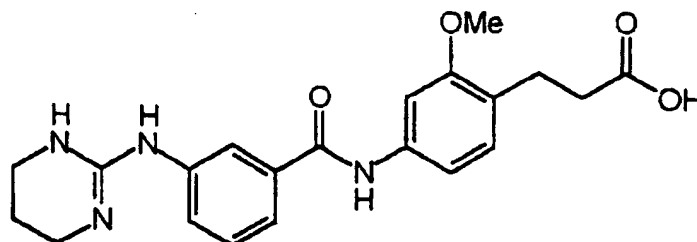
- 251 -

Example 116

2-methoxy-4-[[[3-[(1,4,5,6-tetrahydropyrimidin-2-yl)amino]phenyl]carbonyl]amino]benzenepropanoic acid, trifluoroacetate salt

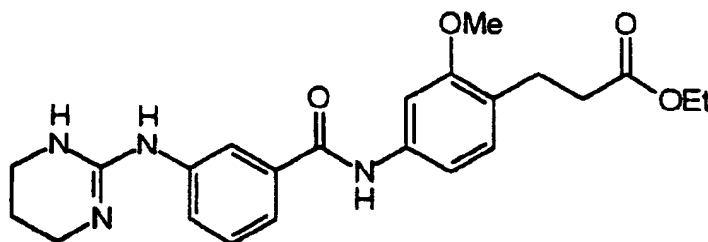
5

10

Step A

15

20



The product from Example 115 Step B (350 mg, 1.56 mmol) and the product of Example 130 (400 mg, 1.56 mmol) were coupled using the procedure of Example 76 Step D to give the desired compound. ^1H NMR was consistent with the proposed structure.

Analysis calculated for $\text{C}_{23}\text{H}_{28}\text{N}_4\text{O}_4 \cdot 1.0 \text{ TFA} \cdot 1.5 \text{ H}_2\text{O}$:

C, 53.09; H, 5.70; N, 9.91.

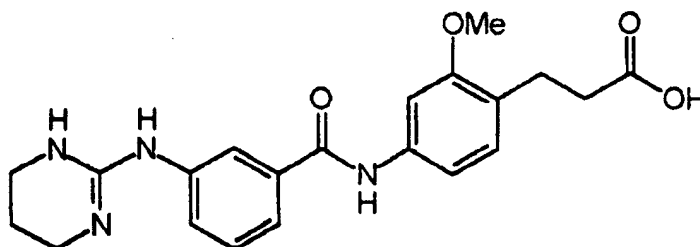
Found: C, 52.94; H, 5.52; N, 9.63.

30

- 252 -

Step B

5



10 The product of Step A was saponified using the conditions of Example 35 to give the above compound.

¹H NMR was consistent with the proposed structure.

Analysis calculated for C₂₃H₂₈N₄O₄•1.0 TFA•0.5 H₂O:

C, 53.18; H, 5.04; N, 10.79.

15

Found: C, 53.46; H, 5.04; N, 10.74.

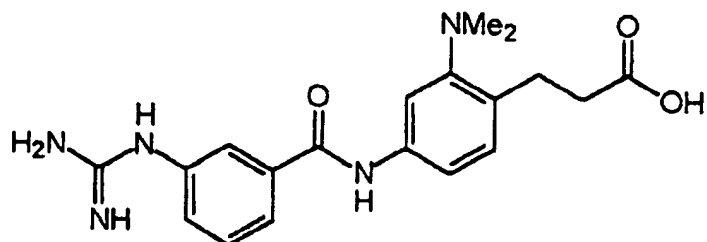
- 253 -

Example 117

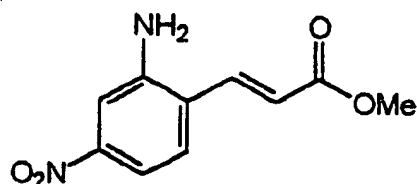
4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-2-(dimethylamino)benzenepropanoic acid,
bis(trifluoroacetate) salt

5

10

StepA

15



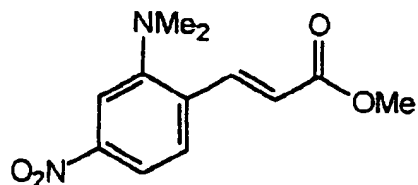
20

2-bromo-5-nitroaniline (2 g, 9.22 mmol) and methyl
acrylate were coupled using the procedure described in
Example 100 Step B to give, after purification by flash
chromatography (20/80 ethyl acetate/hexane), 1.05 g of
product. ¹H NMR was consistent with the proposed
structure.

25

Step B

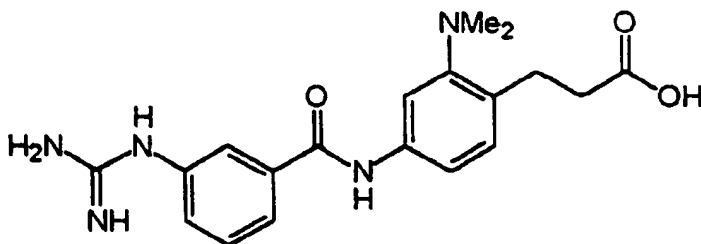
30



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The compound of Step A (500 mg, 2.25 mmol) and potassium carbonate (622 mg, 4.50 mmol) were stirred in of DMF (5 ml) for 1 hour and methyl iodide (5 mL) was added. The mixture was stirred at 50°C overnight, cooled, diluted with water and extracted with ethyl acetate. The solution was dried over MgSO₄ and concentrated to give, after flash chromatography (20/80 ethyl acetate/hexane), 380 mg of product. ¹H NMR was consistent with the proposed structure.

Steps C/D/E



The compound from Step B was reduced, coupled and saponified using the conditions described in Example 115 Steps B and C/D to provide the above compound. ¹H NMR was consistent with the proposed structure.

Analysis calculated for C₁₉H₂₃N₅O₃•2.8 TFA:

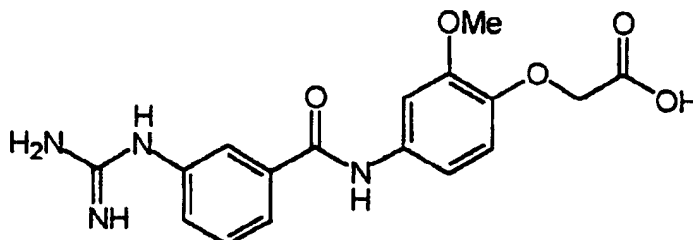
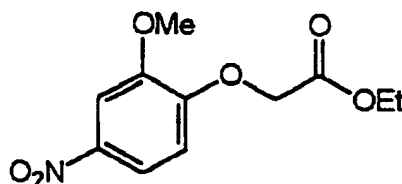
C, 42.90; H, 3.78; N, 10.17.

Found: C, 42.78; H, 3.65; N, 10.03.

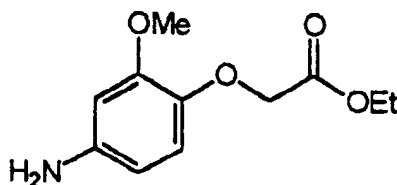
- 255 -

Example 118

[4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-2-methoxyphenoxy]acetic acid,
trifluoroacetate salt

Step A

A solution of 4-nitroguaiacol (2 g, 11.83 mmol) and potassium carbonate (1.96 g, 14.19 mmol) in DMF (20 ml) was stirred at room temperature for 1 hour and ethyl bromoacetate (1.98 g, 11.83 mmol) was added. After stirring at room temperature overnight the reaction mixture was diluted with water and the solid filtered, washed with water and dried to give the product. ¹H NMR was consistent with the proposed structure.

Step B

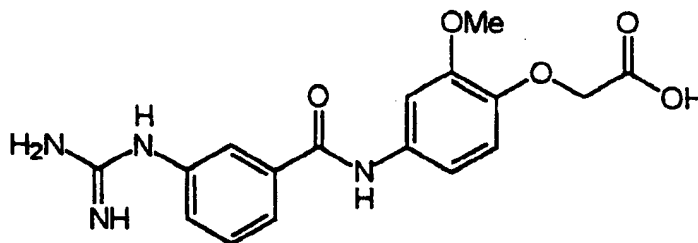
- 256 -

To the compound of Step A (2.9 g, 11.36 mmol) in ethanol (80 ml) was added tin(II) chloride dihydrate (7.69 g, 34.09 mmol) and the reaction mixture stirred at 80°C for 4 hours. The mixture was cooled, neutralized with 10% NaOH and extracted with ethyl acetate. The extract was dried over MgSO₄ and concentrated to give, after flash chromatography (10/90 ethyl acetate/hexane), 660 mg of product. ¹H NMR was consistent with the proposed structure.

10

Steps C/D

15



20

The product of Step B was coupled with 3-guanidinobenzoic acid hydrochloride as described in Example 76 Step D and saponified as described in Example 35 to furnish the above compound. ¹H NMR was consistent with the proposed structure.

25

Analysis calculated for C₁₇H₁₈N₄O₅•1.0 TFA:

C, 48.31; H, 4.05; N, 11.86.

Found:

C, 48.08; H, 3.94; N, 11.69.

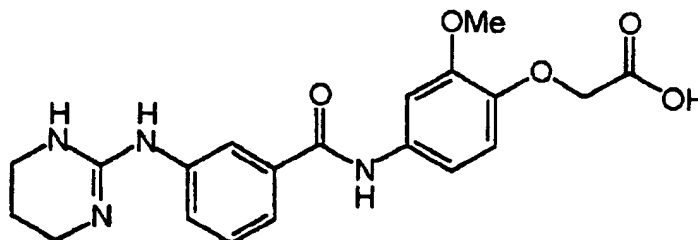
- 257 -

Example 119

[2-methoxy-4-[[[3-[(1,4,5,6-tetrahydropyrimidin-2-yl)-
amino]phenyl]carbonyl]amino]phenoxy]acetic
acid, trifluoroacetate salt

5

10



The above compound was prepared as described for
Example 118, using the product of Example 130 in place of
3-guanidinobenzoic acid hydrochloride. ¹H NMR was
consistent with the proposed structure.

Analysis calculated for C₁₇H₁₈N₄O₅•1.2 TFA•1.0 H₂O:

C, 48.63; H, 4.59; N, 10.13.

Found:

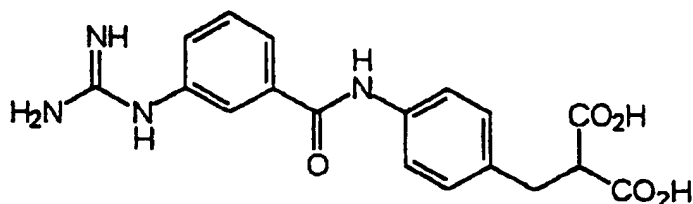
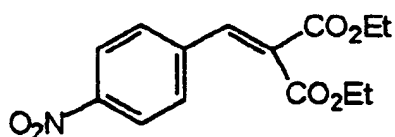
C, 48.23; H, 4.31; N, 9.89.

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Example 120

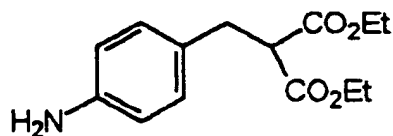
[[4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]phenyl]methyl]-1,3-propanoic acid,
trifluoroacetic salt

5

10 Step A

15

The above compound was prepared using the procedure described in Synthesis-Stuttgart 12, 1026-1027 (1986). ¹H NMR was consistent with the proposed structure.

20 Step B

25

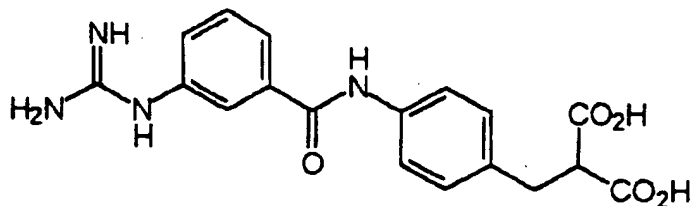
The compound of Step A was reduced using the procedure described in Example 106 Step C to provide the desired aniline. ¹H NMR was consistent with the proposed structure.

30

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Step C

5



The above compound was prepared using the procedures described in Example 127 Step B to Step D/E/F. ¹H NMR was
10 consistent with the proposed structure.

Analysis calculated for C₁₈H₁₆N₄O₅•1.8 TFA• 0.7 H₂O:

C, 44.26; H, 3.30; N, 9.56.

Found:

C, 44.17; H, 2.99; N, 9.75.

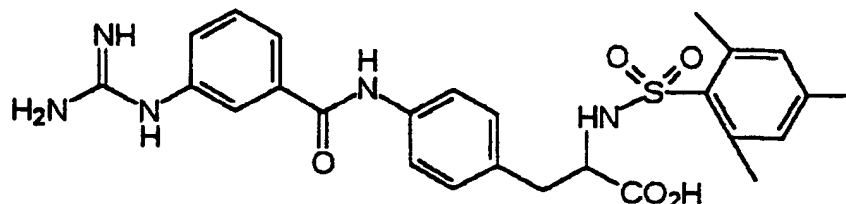
- 260 -

Example 121

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(2,4,6-trimethylphenyl)sulfonyl]-
phenylalanine, trifluoroacetate salt

5

10



The above compound was prepared in the same manner as
described in Example 106, replacing isovaleryl chloride
with 2,4,6-trimethylbenzenesulfonyl chloride. ¹H NMR was
consistent with the proposed structure.

Analysis calculated for C₂₆H₂₉N₅O₅•1.5 TFA• 0.2 H₂O:

C, 49.88; H, 4.46; N, 10.03.

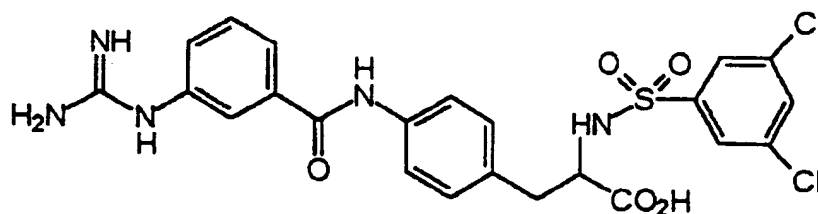
Found:

C, 49.71; H, 4.51; N, 10.05.

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Example 122

[4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(3,5-dichlorophenyl)sulfonyl]phenyl-
alanine, trifluoroacetate salt



The above compound was prepared in the same manner as described in Example 106, replacing isovaleryl chloride with 3,5-dichlobenzenesulfonyl chloride. ^1H NMR was consistent with the proposed structure.

Analysis calculated for $\text{C}_{23}\text{H}_{21}\text{N}_5\text{O}_5\text{SCl}_2 \cdot 1.4 \text{ TFA}$:

C, 51.86; H, 4.85; N, 10.61.

Found:

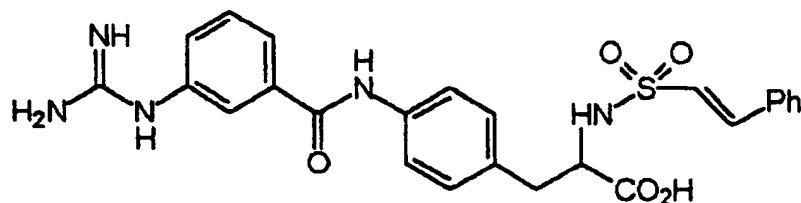
C, 51.57; H, 4.99; N, 11.01.

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Example 123

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(2-phenylethenyl)sulfonyl]phenyl-
alanine, trifluoroacetate salt

5



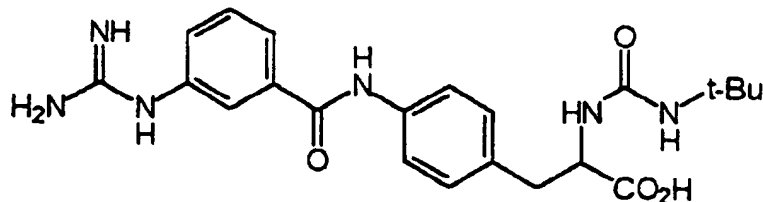
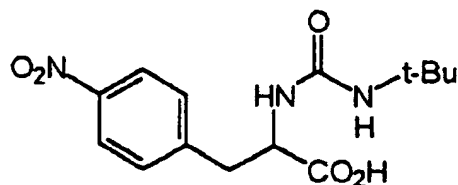
10

The above compound was prepared in the same manner as described in Example 106, replacing isovaleryl chloride with trans- β -styrenesulfonyl chloride. ^1H NMR was consistent with the proposed structure.

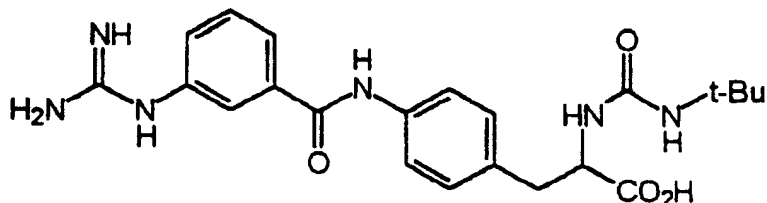
- 263 -

Example 124

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[[[(1,1-dimethylethyl)amino]carbonyl]-
phenylalanine, trifluoroacetate salt

Step A

The product of Example 106 Step A was dissolved in CH_2Cl_2 (8 mL) and cooled to 0°C . t-Butyl isocyanate (139 mg, 1.4 mmol) was added followed by triethylamine (151 mg, 1.5 mmol) and the mixture stirred overnight. The mixture was concentrated in vacuo, the residue dissolved in EtOAc and the solution washed with water and brine, dried over Na_2SO_4 and concentrated to give 0.5 g of a yellow solid. ^1H NMR was consistent with the proposed structure.

Steps B/C/D

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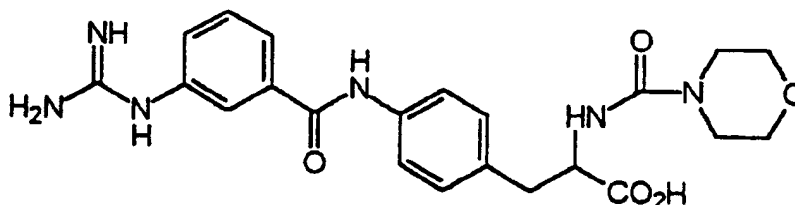
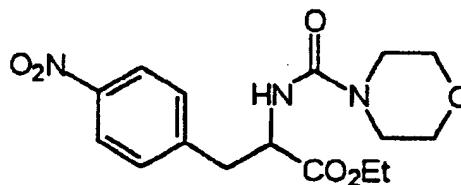
The title compound was prepared from the product of Step A using the procedures described in Example 106 Steps C, D and E. ¹H NMR was consistent with the proposed structure.

5 Analysis calculated for $C_{22}H_{28}N_6O_4 \cdot 1.5$ TFA:
 C, 49.14; H, 4.78; N, 13.75.
 Found: C, 48.99; H, 4.77; N, 14.11.

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Example 125

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(4-morpholinyl)carbonyl]phenylalanine,
bis(trifluoroacetate) salt, monohydrate

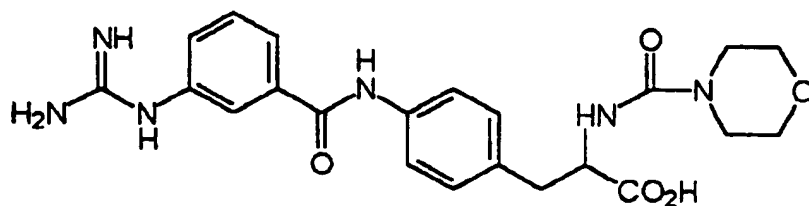
Step A

The compound of Example 106 Step A was dissolved in benzene (15 mL) and cooled to 0°C. Triphosgene (216 mg, 0.73 mmol) was added followed by triethylamine (441 mg, 4.36 mmol) and the mixture refluxed for 2 hours. The mixture was cooled and concentrated in vacuo and the residue dissolved in CH₂Cl₂. Morpholine (190 mg, 2.18 mmol) was added and the mixture stirred at room temperature overnight. The mixture was concentrated in vacuo and the residue dissolved in EtOAc, washed with water and brine and dried over Na₂SO₄. Concentration and purification by flash chromatography (40/60 acetonitrile/toluene) furnished 590 mg of a white solid. ¹H NMR was consistent with the proposed structure.

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Steps B/C/D

5



The title compound was prepared from the product of
Step A using the procedures described in Example 106 Steps
10 C, D and E. ^1H NMR was consistent with the proposed
structure.

Analysis calculated for $\text{C}_{22}\text{H}_{26}\text{N}_6\text{O}_5 \cdot 2.0 \text{ TFA} \cdot 0.1 \text{ H}_2\text{O}$:

C, 45.63; H, 4.15; N, 12.28.

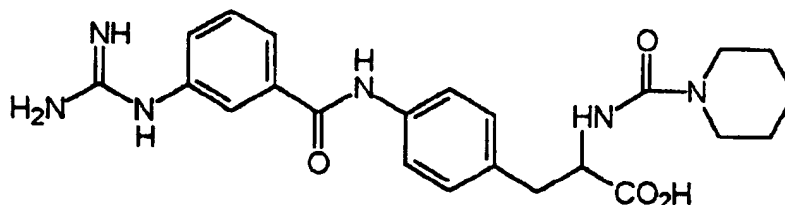
Found:

C, 45.79; H, 4.52; N, 12.12.

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Example 126

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(1-piperidiny)carbonyl]phenyl-
alanine, trifluoroacetate salt



The title compound was prepared in the same manner as described in Example 125, replacing morpholine with piperidine. ^1H NMR was consistent with the proposed structure.

Analysis calculated for $\text{C}_{23}\text{H}_{28}\text{N}_6\text{O}_4 \cdot 1.4 \text{ TFA} \cdot 0.1 \text{ H}_2\text{O}$:

C, 50.47; H, 4.86; N, 13.69.

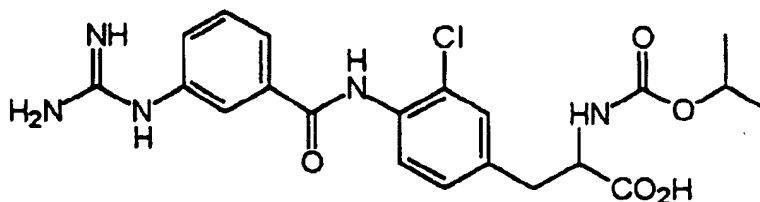
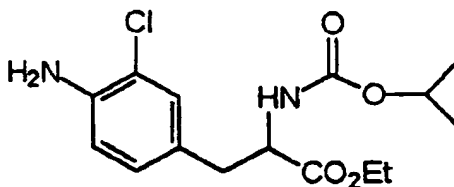
Found:

C, 50.26; H, 4.67; N, 13.85.

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Example 127

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-3-chloro-N-[(1-methylethoxy)carbonyl]-
phenylalanine, trifluoroacetate salt

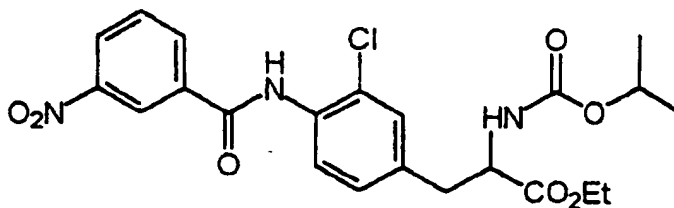
Step A

The compound of Example AE (588 mg, 2.0 mmol) and N-chlorosuccinimide (268 mg, 2.0 mmol) were stirred in acetonitrile (8 mL) at room temperature overnight. The mixture was concentrated in vacuo and the residue suspended in 10% NaHSO₃ and stirred for 2 hours. The mixture was extracted with ethyl acetate (3X) and the organic layers collected, washed with brine, dried over Na₂SO₄ and concentrated. The residue was purified by flash chromatography (99:1 MeOH/CH₂Cl₂) to give 360 mg of the desired compound. ¹H NMR was consistent with the proposed structure.

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Step B

5

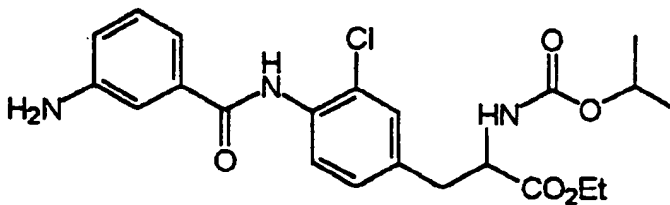


The desired compound was prepared from the product of Step A using the procedure described in Example AQ. ¹H NMR was consistent with the proposed structure.

10

Step C

15

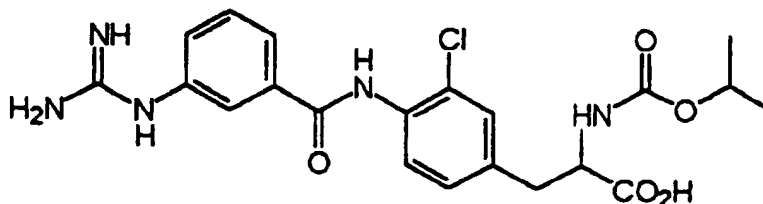


The desired compound was prepared from the product of Step B using the procedure described in Example 106 Step C. ¹H NMR was consistent with the proposed structure.

20

Steps D/E/F

25



The title compound was prepared from the product of Step C using the procedures described in Example 108 Steps A and B/C. ¹H NMR was consistent with the proposed structure.

Analysis calculated for C₂₁H₂₄N₅O₅•1.5 TFA:

C, 45.54; H, 4.06; N, 11.06.

Found:

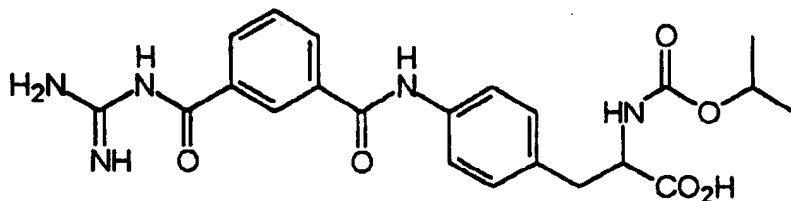
C, 45.46; H, 4.11; N, 10.79.

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Example 128

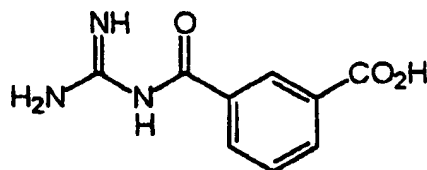
4-[[[3-[[[(aminoiminomethyl) amino] carbonyl] phenyl]-
carbonyl] amino]-N-[(1-methylethoxy) carbonyl]-
phenylalanine, trifluoroacetate salt

5



10

Step A



15

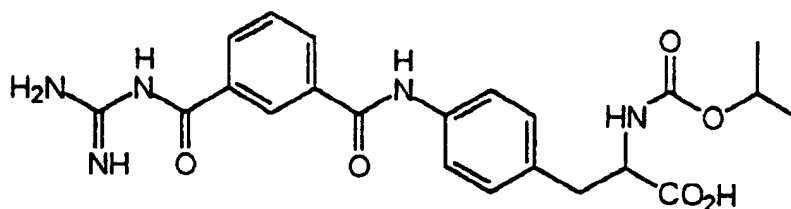
Flask A: A suspension of 60% NaH in mineral oil (1.2 g, 24 mmol washed with hexane before use) in DMF (20 mL) was cooled to 0°C and guanidine hydrochloride (2.4 g, 25 mmol) was added very slowly. The reaction was allowed to stir at room temperature for 2 hours.

Flask B: To a solution of isophthalic acid (0.83 g, 5 mmol) in DMF (5 mL) at 0°C was added 1-methylpiperidine (496 mg, 5 mmol), followed by isobutyl chloroformate (205 mg, 1.5 mmol). The mixture was stirred for 5 minutes and the solution of Flask A was added. The mixture was stirred overnight and concentrated. The residue was purified by reverse-phase HPLC (water/acetonitrile) to afford 300 mg of a white solid. ¹H NMR was consistent with the proposed structure.

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Steps B/C

5



The title compound was prepared from the product of Step A and the compound of Example AE using the procedures described in Example 106 Steps D and E. ¹H NMR was consistent with the proposed structure.

Analysis calculated for C₂₂H₂₅N₅O₆•1.3 TFA:

C, 48.93; H, 4.39; N, 11.60.

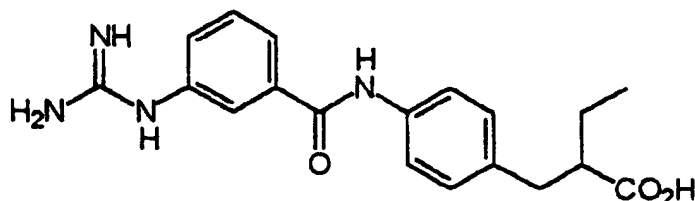
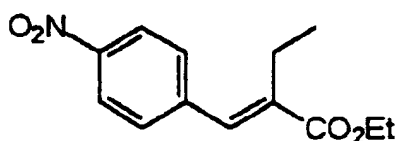
Found: C, 48.74; H, 4.63; N, 11.44.

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Example 129

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]- α -ethylbenzenepropanoic acid,
trifluoroacetate salt

5

10 Step A

15

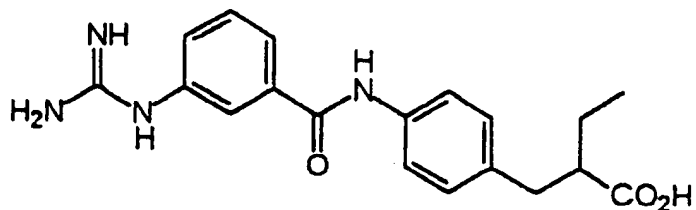
A suspension of 60% NaH in mineral oil (2 g, 51 mmol, washed with hexane before use) in THF (75 mL) was cooled to 0°C and triethyl 2-phosphonobutyrate (7.75 mL, 33 mmol) was added slowly. The white slurry was stirred at 0°C for 1.5 hours and a solution of 4-nitrobenzaldehyde (4.5 g, 30 mmol) in THF (10 mL) was added. The mixture was stirred at room temperature for 4 hours and quenched with water and extracted with EtOAc (3X). The organic layers were collected, washed with brine, dried over Na₂SO₄ and concentrated. The residue was purified by flash chromatography on (4:1 hexane/ethyl acetate) to give 4 g of a yellow solid. ¹H NMR was consistent with the proposed structure.

30

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Steps B/C/D

5



The title compound was prepared from the product of Step A using the procedures described in Example 106 Steps C, D and E. ¹H NMR was consistent with the proposed structure.

Analysis calculated for C₁₉H₂₂N₄O₃•1.2 TFA• 0.5 H₂O:

C, 51.38; H, 4.88; N, 11.20.

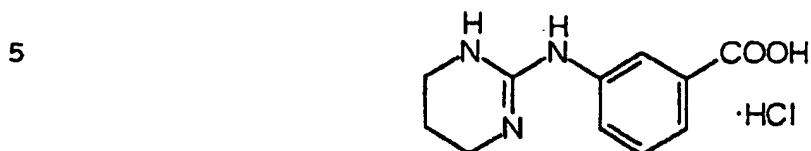
Found:

C, 51.35; H, 4.72; N, 11.36.

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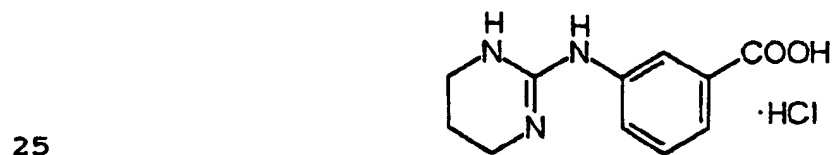
Example 130

3-[(1,4,5,6-tetrahydropyrimidin-2-yl)-
amino]benzoic acid

Step A

15 1-(3-Carboxyphenyl)-2-thiourea (5 g, 0.025 mol)) and
iodomethane (3.62 g, 0.025 mole) in THF (75 mL) were
stirred at reflux for 2 hours. The solvent was removed
and the residue washed with ether (3X) to yield, after
drying under vacuum, the desired salt (7.8 g) as a yellow
solid.

20

Step B

To the product of Step A (10.1 g, 0.03 mol) in DMF
(15 mL) was added 1,3-diaminopropane (2.22 g, 0.03 mol),
triethylamine (3.9 g, 0.03 mol), and DMAP (420 mg). The
30 reaction mixture was heated at 140-150°C for 4.5 hours and
cooled to room temperature. H₂O (30 mL) was added and,
after stirring for 15 minutes, the precipitate filtered
and washed with H₂O. The precipitate was slurried in H₂O
and made acidic with concentrated HCl. The solution was

- 275 -

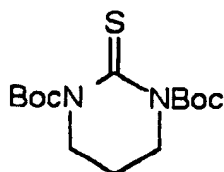
lyophilized and the residue washed 2X with isopropyl ether and dried, furnishing the above compound (4.0 g) as a white solid. ^1H NMR and MS were consistent with the desired structure.

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Example 131

N,N' -bis-(Boc)-2-(1H)-tetrahydropyrimidinethione

5

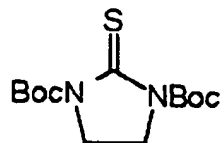


The above compound was prepared from 2-(1H)-
tetrahydropyrimidinethione using the procedure described
10 in Example AZ. ¹H NMR was consistent with the desired
structure.

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Example 132

N,N' -bis-(Boc) -2-imidazolidinethione



5

The above compound was prepared from 2-
imidazolidinethione using the procedure described in
10 Example A2. ¹H NMR was consistent with the desired
structure.

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The activity of the compounds of the present invention was tested in the following assays. The results of testing in the assays are tabulated in Table 1.

5 VITRONECTIN ADHESION ASSAY

MATERIALS

 Human vitronectin receptor($\alpha_v\beta_3$) was purified from human placenta as previously described [Pytela et al., Methods in Enzymology, 144:475-489 (1987)]. Human
10 vitronectin was purified from fresh frozen plasma as previously described [Yatohgo et al., Cell Structure and Function, 13:281-292 (1988)]. Biotinylated human vitronectin was prepared by coupling NHS-biotin from Pierce Chemical Company (Rockford, IL) to purified
15 vitronectin as previously described [Charo et al., J. Biol. Chem., 266(3):1415-1421 (1991)]. Assay buffer, OPD substrate tablets, and RIA grade BSA were obtained from Sigma (St. Louis, MO). Anti-biotin antibody was obtained from Calbiochem (La Jolla, CA). Linbro
20 microtiter plates were obtained from Flow Labs (McLean, VA). ADP reagent was obtained from Sigma (St. Louis, MO).

METHODS

25 Solid Phase Receptor Assays

 This assay was essentially the same as previously reported [Niiya et al., Blood, 70:475-483 (1987)]. The purified human vitronectin receptor ($\alpha_v\beta_3$) was diluted from stock solutions to 1.0 $\mu\text{g/mL}$ in Tris-buffered saline
30 containing 1.0 mM Ca^{++} , Mg^{++} , and Mn^{++} , pH 7.4 (TBS^{+++}). The diluted receptor was immediately transferred to Linbro microtiter plates at 100 $\mu\text{L/well}$ (100 ng receptor/well). The plates were sealed and incubated overnight at 4°C to allow the receptor to bind to the wells. All remaining

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steps were at room temperature. The assay plates were emptied and 200 μ L of 1% RIA grade BSA in TBS⁺⁺⁺ (TBS⁺⁺⁺/BSA) were added to block exposed plastic surfaces. Following a 2 hour incubation, the assay plates were

5 washed with TBS⁺⁺⁺ using a 96 well plate washer. Logarithmic serial dilution of the test compound and controls were made starting at a stock concentration of 2 mM and using 2 nM biotinylated vitronectin in TBS⁺⁺⁺/BSA as the diluent. This premixing of labeled ligand with

10 test (or control) ligand, and subsequent transfer of 50 μ L aliquots to the assay plate was carried out with a CETUS Propette robot; the final concentration of the labeled ligand was 1 nM and the highest concentration of test compound was 1.0×10^{-4} M. The competition occurred for

15 two hours after which all wells were washed with a plate washer as before. Affinity purified horseradish peroxidase labeled goat anti-biotin antibody was diluted 1:3000 in TBS⁺⁺⁺/BSA and 125 μ L were added to each well. After 30 minutes, the plates were washed and incubated

20 with OPD/H₂O₂ substrate in 100 mM/L Citrate buffer, pH 5.0. The plate was read with a microtiter plate reader at a wavelength of 450 nm and when the maximum-binding control wells reached an absorbance of about 1.0, the final A₄₅₀ were recorded for analysis. The data were

25 analyzed using a macro written for use with the EXCEL[™] spreadsheet program. The mean, standard deviation, and %CV were determined for duplicate concentrations. The mean A₄₅₀ values were normalized to the mean of four maximum-binding controls (no competitor added) (B-MAX). The

30 normalized values were subjected to a four parameter curve fit algorithm [Rodbard et al., Int. Atomic Energy Agency, Vienna, pp 469 (1977)], plotted on a semi-log scale, and the computed concentration corresponding to inhibition of 50% of the maximum binding of biotinylated vitronectin

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(IC₅₀) and corresponding R² was reported for those compounds exhibiting greater than 50% inhibition at the highest concentration tested; otherwise the IC₅₀ is reported as being greater than the highest concentration tested. β -[[2-[[5-[(aminoiminomethyl)amino]-1-oxopentyl]amino]-1-oxoethyl]amino]-3-pyridinepropanoic acid [USSN 08/375,338, Example 1] which is a potent $\alpha_v\beta_3$ antagonist (IC₅₀ in the range 3-10 nM) was included on each plate as a positive control.

10

PURIFIED IIb/IIIa RECEPTOR ASSAY

MATERIALS

Human fibrinogen receptor ($\alpha_{IIb}\beta_3$) was purified from outdated platelets. (Pytela, R., Pierschbacher, M.D., Argraves, S., Suzuki, S., and Rouslahti, E. "Arginine-Glycine-Aspartic acid adhesion receptors", Methods in Enzymology 144(1987):475-489.) Human vitronectin was purified from fresh frozen plasma as described in Yatohgo, T., Izumi, M., Kashiwagi, H., and Hayashi, M., "Novel purification of vitronectin from human plasma by heparin affinity chromatography," Cell Structure and Function 13(1988):281-292. Biotinylated human vitronectin was prepared by coupling NHS-biotin from Pierce Chemical Company (Rockford, IL) to purified vitronectin as previously described. (Charo, I.F., Nannizzi, L., Phillips, D.R., Hsu, M.A., Scarborough, R.M., "Inhibition of fibrinogen binding to GP IIb/IIIa by a GP IIIa peptide", J. Biol. Chem. 266(3)(1991): 1415-1421.) Assay buffer, OPD substrate tablets, and RIA grade BSA were obtained from Sigma (St. Louis, MO). Anti-biotin antibody was obtained from Calbiochem (La Jolla, CA). Linbro microtiter plates were obtained from Flow Labs (McLean, VA). ADP reagent was obtained from Sigma (St. Louis, MO).

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METHODSSolid Phase Receptor Assays

This assay is essentially the same reported in Niiya, K., Hodson, E., Bader, R., Byers-Ward, V. Koziol, J.A., Plow, E.F. and Ruggeri, Z.M., "Increased surface expression of the membrane glycoprotein IIb/IIIa complex induced by platelet activation: Relationships to the binding of fibrinogen and platelet aggregation", Blood 70(1987):475-483. The purified human fibrinogen receptor ($\alpha_{IIb}\beta_3$) was diluted from stock solutions to 1.0 $\mu\text{g/mL}$ in Tris-buffered saline containing 1.0 mM Ca^{++} , Mg^{++} , and Mn^{++} , pH 7.4 (TBS^{+++}). The diluted receptor was immediately transferred to Linbro microtiter plates at 100 $\mu\text{L/well}$ (100 ng receptor/well). The plates were sealed and incubated overnight at 4°C to allow the receptor to bind to the wells. All remaining steps were at room temperature. The assay plates were emptied and 200 μL of 1% RIA grade BSA in TBS^{+++} ($\text{TBS}^{+++}/\text{BSA}$) were added to block exposed plastic surfaces. Following a 2 hour incubation, the assay plates were washed with TBS^{+++} using a 96 well plate washer. Logarithmic serial dilution of the test compound and controls were made starting at a stock concentration of 2 mM and using 2 nM biotinylated vitronectin in $\text{TBS}^{+++}/\text{BSA}$ as the diluent. This premixing of labeled ligand with test (or control) ligand, and subsequent transfer of 50 μL aliquots to the assay plate was carried out with a CETUS Propette robot; the final concentration of the labeled ligand was 1 nM and the highest concentration of test compound was 1.0×10^{-4} M. The competition occurred for two hours after which all wells were washed with a plate washer as before. Affinity purified horseradish peroxidase labeled goat anti-biotin antibody was diluted 1:3000 in $\text{TBS}^{+++}/\text{BSA}$ and 125 μL were

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added to each well. After 30 minutes, the plates were washed and incubated with ODD/H₂O₂ substrate in 100 mM/L citrate buffer, pH 5.0. The plate was read with a microtiter plate reader at a wavelength of 450 nm and when
5 the maximum-binding control wells reached an absorbance of about 1.0, the final A₄₅₀ were recorded for analysis. The data were analyzed using a macro written for use with the EXCEL[™] spreadsheet program. The mean, standard deviation, and %CV were determined for duplicate concentrations. The
10 mean A₄₅₀ values were normalized to the mean of four maximum-binding controls (no competitor added) (B-MAX). The normalized values were subjected to a four parameter curve fit algorithm, [Robard et al., Int. Atomic Energy Agency, Vienna, pp 469 (1977)], plotted on a semi-log
15 scale, and the computed concentration corresponding to inhibition of 50% of the maximum binding of biotinylated vitronectin (IC₅₀) and corresponding R² was reported for those compounds exhibiting greater than 50% inhibition at the highest concentration tested; otherwise the IC₅₀ is
20 reported as being greater than the highest concentration tested. β -[[2-[[5-[(aminoiminomethyl)amino]-1-oxopentyl]amino]-1-oxoethyl]amino]-3-pyridinepropanoic acid [USSN 08/375,338, Example 1] which is a potent $\alpha_v\beta_3$ antagonist (IC₅₀ in the range 3-10 nM) was included on
25 each plate as a positive control.

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TABLE I

	Example	AvB3 IC50 (nM)	I Ib/IIIa IC50 (nM)
5	2	0.18	8.08
	6	0.51	49.5
	10	1.17	26.9
	11	2.2	4
10	12	>100000	69200
	19	>100000	1780
	20	>100000	39900
	24	1220	4530
	28	555	10100
15	29	4000	1280
	30	6370	3260
	31	114	66.5
	33	73.6	7.37
	34	45.2	2200
20	35	2.69	148
	36	1010	10500
	37	97.8	2380
	38	106	3360
	40	557	748
25	41	1080	5700
	44	120	4310
	45	39.4	603

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	Example	AvB3 IC50 (nM)	IIb/IIIa IC50 (nM)
5	47	5.89	51.1
	48	0.63	15.6
	49	1.1	31.4
	50	0.8	21.7
	51	0.63	0.71
10	52	0.33	1.88
	53	0.56	15.3
	54	0.77	36.5
	55	0.51	14.3
	56	0.43	6.35
15	57	14.5	318
	58	48400	57200
	59	4480	478
	60	3370	3810
	61	470	666
20	62	28100	14800
	64	21.2	93.1
	65	4760	21900
	68	782	103
	69	94	37
	71	38.7	8.3
	73	141	40
	75	14.5	330
	77	1.6	1260

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	Example	AvB3 IC50 (nM)	IIb/IIIa IC50 (nM)
	78	5980	397
	79	2290	10200
	80	44200	19800
	81	19.1	129
5	82	264	332
	83	1900	474
	84	16300	486
	85	827	12600
	85C	39600	66000
10	86	307	3140
	86C	56800	>100000
	87	0.44	85.3
	89	2070	16000
	90	23700	34500
15	100	6	44
	101A	9	57
	101B	35	286
	102	6	4710
	103	269	3660
20	104	506	1600
	105	2	267
	106	5	77
	107	0.5	1570
	108	4	109

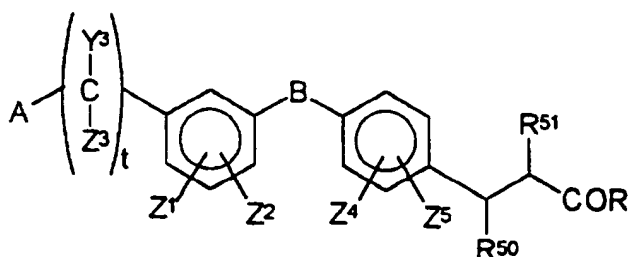
- 286 -

	Example	AvB3 IC50 (nM)	IIb/IIIa IC50 (nM)
5	109	5	111
	110	4	326
	111	0.1	6
	112	0.3	13
	113	1	253
10	114	0.2	1000
	115	24	173
	116	6	3180
	117	129	452
	118	3	54
15	119	0.4	772
	120	14	384
	121	1	2
	122	7	50
	123	0.5	1.5
20	124	3	285
	125	7	260
	126	2	36
	127	0.6	22
	128	216	20
	129	265	5260

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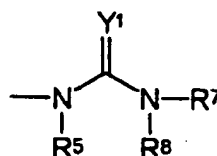
What is claimed is:

1. A compound of the formula



or a pharmaceutically acceptable salt thereof,
wherein

A is



wherein Y¹ is selected from the group consisting of N-R², O, and S;

R² is selected from the group consisting of H; alkyl; aryl; hydroxy; alkoxy; cyano; nitro; amino; aminocarbonyl; alkenyl; alkynyl; alkyl optionally substituted with one or more substituent selected from lower alkyl, halogen, hydroxyl, haloalkyl, cyano, nitro, carboxyl, amino, alkoxy, aryl or aryl optionally substituted with one or more halogen, haloalkyl, lower alkyl, alkoxy, cyano, alkylsulfonyl, alkylthio, nitro, carboxyl, amino, hydroxyl, sulfonic acid, sulfonamide, aryl, fused aryl, monocyclic heterocycles, or fused monocyclic heterocycles; aryl optionally substituted with one or more substituent selected from halogen, haloalkyl, hydroxy, lower alkyl, alkoxy, methylenedioxy, ethylenedioxy, cyano,

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nitro, alkylthio, alkylsulfonyl, sulfonic acid, sulfonamide, carboxyl derivatives, amino, aryl, fused aryl, monocyclic heterocycles and fused monocyclic heterocycle; monocyclic heterocycles; and monocyclic heterocycles optionally substituted with one or more substituent selected from halogen, haloalkyl, lower alkyl, alkoxy, amino, nitro, hydroxy, carboxyl derivatives, cyano, alkylthio, alkylsulfonyl, sulfonic acid, sulfonamide, aryl or fused aryl; or

R^2 taken together with R^7 forms a 4-12 membered dinitrogen containing heterocycle optionally substituted with one or more substituent selected from the group consisting of lower alkyl, hydroxy and phenyl and wherein said ring optionally contains a heteroatom selected from the group consisting of O and S;

or R^2 taken together with R^7 forms a 5 membered heteroaromatic ring optionally substituted with amino;

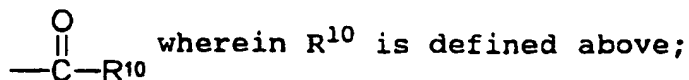
or R^2 taken together with R^7 forms a 5 membered heteroaromatic ring fused with a phenyl group;

R^7 (when not taken together with R^2) and R^8 are independently selected from the group consisting of H; alkyl; alkenyl; alkynyl; aralkyl; cycloalkyl; bicycloalkyl; aryl; acyl; benzoyl; alkyl optionally substituted with one or more substituent selected from lower alkyl, halogen, hydroxy, haloalkyl, cyano, nitro, carboxyl derivatives, amino, alkoxy, thio, alkylthio, sulfonyl, aryl, aralkyl, aryl optionally substituted with one or more substituent selected

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from halogen, haloalkyl, lower alkyl, alkoxy, methylenedioxy, ethylenedioxy, alkylthio, haloalkylthio, thio, hydroxy, cyano, nitro, carboxyl derivatives, aryloxy, amido, acylamino, amino, alkylamino, dialkylamino, trifluoroalkoxy, trifluoromethyl, sulfonyl, alkylsulfonyl, haloalkylsulfonyl, sulfonic acid, sulfonamide, aryl, fused aryl, monocyclic heterocycles, fused monocyclic heterocycles; aryl optionally substituted with one or more substituent selected from halogen, haloalkyl, lower alkyl, alkoxy, methylenedioxy, ethylenedioxy, alkylthio, haloalkylthio, thio, hydroxy, cyano, nitro, carboxyl derivatives, aryloxy, amido, acylamino, amino, alkylamino, dialkylamino, trifluoroalkoxy, trifluoromethylsulfonyl, alkylsulfonyl, sulfonic acid, sulfonamide, aryl, fused aryl, monocyclic heterocycles, or fused monocyclic heterocycles; monocyclic heterocycles; monocyclic heterocycles optionally substituted with one or more substituent selected from halogen, haloalkyl, lower alkyl, alkoxy, aryloxy, amino, nitro, hydroxy, carboxyl derivatives, cyano, alkylthio, alkylsulfonyl, aryl, fused aryl; monocyclic and bicyclic heterocyclicalkyls; $-SO_2R^{10}$ wherein R^{10} is selected from the group consisting of alkyl, aryl and monocyclic heterocycles, all optionally substituted with one or more substituent selected from the group consisting of halogen, haloalkyl, alkyl, alkoxy, cyano, nitro, amino, acylamino, trifluoroalkyl, amido, alkylaminosulfonyl, alkylsulfonyl, alkylsulfonylamino, alkylamino, dialkylamino, trifluoromethylthio, trifluoroalkoxy, trifluoromethylsulfonyl, aryl, aryloxy, thio, alkylthio, and monocyclic heterocycles; and

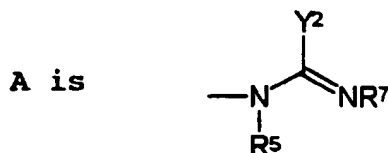
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or NR^7 and R^8 taken together form a 4-12 membered mononitrogen containing monocyclic or bicyclic ring optionally substituted with one or more substituent selected from lower alkyl, carboxyl derivatives, aryl or hydroxy and wherein said ring optionally contains a heteroatom selected from the group consisting of O, N and S;

R^5 is selected from the group consisting of H, alkyl, alkenyl, alkynyl, benzyl, and phenethyl;

or



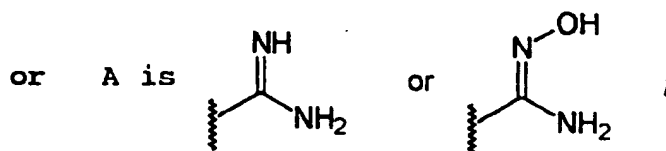
wherein Y^2 is selected from the group consisting of H, alkyl; cycloalkyl; bicycloalkyl; aryl; monocyclic heterocycles; alkyl optionally substituted with aryl which can also be optionally substituted with one or more substituent selected from halo, haloalkyl, alkyl, nitro, hydroxy, alkoxy, aryloxy, aryl, or fused aryl; aryl optionally substituted with one or more substituent selected from halo, haloalkyl, hydroxy, alkoxy, aryloxy, aryl, fused aryl, nitro, methylenedioxy, ethylenedioxy, or alkyl; alkynyl; alkenyl; $-\text{S}-\text{R}^9$ and $-\text{O}-\text{R}^9$ wherein R^9 is selected from the group consisting of H; alkyl; aralkyl; aryl; alkenyl; and alkynyl; or R^9 taken together with R^7

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forms a 4-12 membered mononitrogen containing sulfur or oxygen containing heterocyclic ring; and

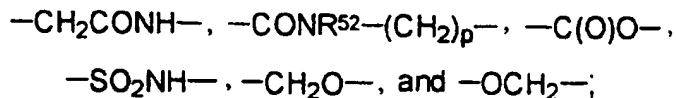
R^5 and R^7 are as defined above;

or γ^2 (when γ^2 is carbon) taken together with R^7 forms a 4-12 membered mononitrogen containing ring optionally substituted with alkyl, aryl or hydroxy;



Z^1 , Z^2 , Z^4 and Z^5 are independently selected from the group consisting of H; alkyl; hydroxy; alkoxy; aryloxy; aralkoxy; halogen; haloalkyl; haloalkoxy; nitro; amino; aminoalkyl; alkylamino; dialkylamino; cyano; alkylthio; alkylsulfonyl; carboxyl derivatives; carboxyalkyl; alkoxy-carbonylalkyl; acetamide; aryl; fused aryl; cycloalkyl; thio; monocyclic heterocycles; fused monocyclic heterocycles; and A, wherein A is defined above;

B is selected from the group consisting of



wherein p is an integer selected from the group consisting of 0, 1 and 2;

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wherein R^{50} is selected from the group consisting of H, alkyl, aryl, carboxyl derivative and -
 $\text{CONHCH}_2\text{CO}_2R^{53}$ wherein R^{53} is H or lower alkyl;

R^{52} is selected from the group consisting of H or alkyl;

R^{51} is selected from the group consisting of H, alkyl, carboxyl derivatives,

$-\text{NCO}_2R^{54}$, $-\text{NHSO}_2R^{54}$, $-\text{NHCONHR}^{54}$,
H

$-\text{NHCOR}^{54}$, $-\text{NHCOR}^{55}$ and amino;

wherein R^{54} is selected from the group consisting of H, alkyl, cycloalkyl, aryl, aralkyl, aralkenyl and aryl substituted by one or more alkyl or halo;
wherein R^{55} is selected from the group consisting of N-substituted pyrrolidinyl, piperidinyl and morpholinyl;

t is an integer 0, 1 or 2;

R is $X-R^3$ wherein X is selected from the group consisting of O, S and NR^4 , wherein R^3 and R^4 are independently selected from the group consisting of hydrogen; alkyl; alkenyl; alkynyl; haloalkyl; aryl; arylalkyl; sugars; steroids and in the case of the free acid, all pharmaceutically acceptable salts thereof; and

Y^3 and Z^3 are independently selected from the group consisting of H, alkyl, aryl, cycloalkyl and aralkyl or Y^3 and Z^3 taken together with C form a carbonyl.

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2. A compound according to Claim 1 wherein t is 0.
3. A compound according to Claim 2 wherein B is -CONR⁵²-.
4. A compound according to Claim 3 selected from the group consisting of

4-[[[3-[(aminoiminomethyl)amino]-5-(trifluoromethyl)-phenyl]carbonyl]amino]-N-[(2-methylpropoxy)-carbonyl]-L-phenylalanine, methyl ester;

4-[[[3-[(aminoiminomethyl)amino]-5-(trifluoromethyl)phenyl]carbonyl]amino]-N-[(2-methylpropoxy)carbonyl]phenylalanine, methyl ester;

N-[(2-methylpropoxy)carbonyl]-4-[[[3-[(3,4,5,6-tetrahydro-2H-azepin-7-yl)amino]-5-(trifluoromethyl)phenyl]carbonyl]amino]-phenylalanine, methyl ester;

N-[(2-methylpropoxy)carbonyl]-4-[[[3-[(3,4,5,6-tetrahydro-2H-azepin-7-yl)amino]-5-(trifluoromethyl)-phenyl]carbonyl]amino]phenylalanine;

N-(butylsulfonyl)-4-[[[3-[(3,4,5,6-tetrahydro-2H-azepin-7-yl)amino]-5-(trifluoromethyl)phenyl]-carbonyl]amino]phenylalanine, methyl ester;

N-(butylsulfonyl)-4-[[[3-[(3,4,5,6-tetrahydro-2H-azepin-7-yl)amino]-5-(trifluoromethyl)phenyl]-carbonyl]amino]phenylalanine;

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4-[[[3-[(aminoiminomethyl)amino]-
5-(trifluoromethyl)phenyl]carbonyl]-
amino]-N-(butylsulfonyl)-phenylalanine,
methyl ester;

4-[[[3-[(aminoiminomethyl)amino]-
5-(trifluoromethyl)phenyl]carbonyl]-
amino]-N-(butylsulfonyl)phenylalanine;

N-(butylsulfonyl)-4-[[[3-[(3,4,5,6-tetrahydro-
2H-azepin-7-yl)amino]phenyl]carbonyl]amino]-
phenylalanine, methyl ester;

N-(butylsulfonyl)-4-[[[3-[(3,4,5,6-
tetrahydro-2H-azepin-7-yl)amino]phenyl]-
carbonyl]amino]phenylalanine;

ethyl 4-[[[3-[amino(hydroxyimino)methyl]-
phenyl]carbonyl]amino]benzenepropanoate;

ethyl 3-[[[3-(aminoiminomethyl)phenyl]-
carbonyl]amino]methyl]benzenepropanoate;

ethyl 3-[[[3-(aminoiminomethyl)phenyl]-
carbonyl]amino]benzenepropanoate;

4-[[[3-(aminoiminomethyl)phenyl]carbonyl]-
amino]benzenepropanoic acid;

ethyl 4-[[[3-[(aminocarbonyl)amino]phenyl]-
carbonyl]amino]benzenepropanoate;

4-[[[3-[(aminocarbonyl)amino]phenyl]-
carbonyl]amino]benzenepropanoic acid;

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ethyl 4-[[[3-[[[(phenylmethyl)amino]carbonyl]-amino]phenyl]carbonyl]amino]benzenepropanoate;

4-[[[3-[[[(phenylmethyl)amino]carbonyl]amino]-phenyl]carbonyl]amino]benzenepropanoic acid;

1,1-dimethylethyl 4-[[[3-[(aminoiminomethyl)-amino]phenyl]carbonyl]amino]- β -methylbenzenepropanoate;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-amino]- β -methylbenzenepropanoic acid;

1,1-dimethylethyl 4-[[[3-[(aminoiminomethyl)-amino]phenyl]carbonyl]amino]- β -phenyl benzenepropanoate;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-amino]- β -phenylbenzenepropanoic acid;

N-acetyl-4-[[[3-[(aminoiminomethyl)amino]-phenyl]carbonyl]amino]phenylalanine ethyl ester;

N-acetyl-4-[[[3-[(aminoiminomethyl)amino]-phenyl]carbonyl]amino]phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-amino]-N-(phenylcarbonyl)phenylalanine ethyl ester;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-amino]-N-(methylsulfonyl)phenylalanine ethyl ester;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-amino]-N-(ethoxycarbonyl)phenylalanine ethyl ester;

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4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(1-methylethoxy)carbonyl]-
phenylalanine ethyl ester;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-(phenylsulfonyl)phenylalanine ethyl ester;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-(butylsulfonyl)phenylalanine ethyl ester;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(2-methylpropoxy)carbonyl]-
phenylalanine, ethyl ester;

4-[[[3-[(Aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(phenoxy)carbonyl]phenylalanine,
ethyl ester;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(1,1-dimethylethoxy)carbonyl]-
phenylalanine ethyl ester;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(1,1-dimethylethoxy)carbonyl]-
phenylalanine, methyl ester;

4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]amino]-N-[(1,1-dimethylethoxy)-
carbonyl]-L-phenylalanine, ethyl ester;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-(phenylcarbonyl)phenylalanine;

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4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-(methylsulfonyl)phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-(ethoxycarbonyl)phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(1-methylethoxy)carbonyl]phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-(phenylsulfonyl)phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-(butylsulfonyl)phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(2-methylpropoxy)carbonyl]phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-(methoxycarbonyl)phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]amino]-N-[(1,1-dimethylethoxy)-
carbonyl]-phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(1,1-dimethylethoxy)carbonyl]-
L-phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(1,1-dimethylethoxy)carbonyl]-
phenylalanine;

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ethyl 4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]methylamino]benzenepropanoate;

4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]methylamino]benzenepropanoic acid;

ethyl 4-[[[3-[(aminoiminomethyl)amino]-
phenyl]carbonyl]amino]benzenepropanoate;

4-[[[3-[(aminoiminomethyl)amino]phenyl]
carbonyl]amino]benzenepropanoic acid;

4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]amino]phenylalanine;

[4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]amino]phenyl]butanedioic acid;

4-(1,1-dimethylethyl) 1-ethyl[4-[[[3-
[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]phenyl]butanedioate;

1-ethyl hydrogen[4-[[[3-[(aminoiminomethyl)amino]-
phenyl]carbonyl]amino]-phenyl]butanedioate;

1,1-dimethylethyl 4-[[[3-[(aminoiminomethyl)amino]-
phenyl]carbonyl]-amino]- β -[[[(ethoxycarbonyl)-
methyl]amino]-carbonyl]benzenepropanoate;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-beta-[(carboxymethyl)amino]-
carbonyl]-benzenepropanoic acid;

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1,1-dimethylethyl 4-[[[3-[(aminoiminomethyl)amino]-phenyl]carbonyl]amino]-3-hydroxybenzenepropanoate;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-amino]-3-hydroxybenzenepropanoic acid;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-amino]-N-[(1,1-dimethylethoxy)carbonyl]-2-methoxyphenylalanine, ethyl ester;

4-[[[3-[(4,5-dihydro-4-oxo-1H-imidazol-2-yl)amino]-phenyl]carbonyl]amino]-N-[(1-methylethoxy)-carbonyl]phenylalanine;

4-[[[3-[[[(cyanoimino)[(phenylmethyl)-amino]methyl]amino]phenyl]carbonyl]amino]-benzenepropanoic acid;

[4-[2-[[3-[(aminoiminomethyl)amino]phenyl]-amino]-2-oxoethyl]phenoxy]acetic acid;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-carbonyl]amino]benzene-1,2-dipropanoic acid;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-amino]-2-carboxybenzenepropanoic acid;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-amino]-2-methoxycarbonyl]benzenepropanoic acid;

2-carboxy-4-[[[3-[(1,4,5,6-tetrahydropyrimidin-2-yl)amino]phenyl]carbonyl]amino]-benzenepropanoic acid;

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(±) 4-[[[3-[3-amino-1H-1,2,4-triazol-5-yl)amino]-phenyl]carbonyl]amino]-N-[(1-methylethoxy)carbonyl]phenylalanine;

(±) 4-[[[3-[(5-amino-1,2,4-oxadiazol-3-yl)amino]-phenyl]carbonyl]amino]-N-[(1-methylethoxy)carbonyl]phenylalanine;

4-[[[3-[[amino[(aminocarbonyl)imino]methyl]-amino]phenyl]carbonyl]amino]-N-[(1-methylethoxy)carbonyl]phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-amino]-N-[(3-methyl-1-oxobutyl)phenylalanine;

N-(3-methyl-1-oxobutyl)-4-[[[3-[(1,4,5,6-tetrahydropyrimidin-2-yl)amino]-5-(trifluoromethyl)phenyl]-carbonyl]amino]phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]-5-(trifluoromethyl)phenyl]carbonyl]amino]-N-(3-methyl-1-oxobutyl)phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-amino]-N-(3,3-dimethyl-1-oxobutyl)phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]-5-(trifluoromethyl)-phenyl]carbonyl]amino]-N-(3,3-dimethyl-1-oxobutyl)phenylalanine;

N-(phenylsulfonyl)-4-[[[3-[(1,4,5,6-tetrahydropyrimidin-2-yl)amino]phenyl]carbonyl]-amino]phenylalanine;

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4-[[[3-[(4,5-dihydro-1H-imidazol-2-yl)amino]phenyl]-
carbonyl]amino-N-(phenylsulfonyl)phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]amino]-
2-methoxy-N-[(1-methylethoxy)carbonyl]phenylalanine;

2-methoxy-N-[(1-methylethoxy)carbonyl]-4-
[[[3-[(1,4,5,6-tetrahydropyrimidin-2-yl)amino]-
phenyl]carbonyl]amino]phenylalanine;

4-[[[3-(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-2-methoxybenzenepropanoic acid;

2-methoxy-4-[[[3-[(1,4,5,6-tetrahydropyrimidin-2-yl)-
amino]phenyl]carbonyl]amino]benzenepropanoic acid;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-2-(dimethylamino)benzenepropanoic acid;

[4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-2-methoxyphenoxy]acetic acid;

[2-methoxy-4-[[[3-[(1,4,5,6-tetrahydropyrimidin-2-yl)-
amino]phenyl]carbonyl]amino]phenoxy]acetic acid;

[[4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]phenyl]methyl]-1,3-propanoic acid;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(2,4,6-trimethylphenyl)-
sulfonyl]phenylalanine;

[4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]amino]-
N-[(3,5-dichlorophenyl)sulfonyl]phenylalanine;

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4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]amino]-
N-[(2-phenylethenyl)sulfonyl]phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[[[1,1-dimethylethyl)amino]-
carbonyl]phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(4-morpholinyl)carbonyl]phenylalanine;

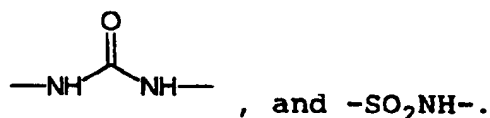
4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(1-pipridinyl)carbonyl]phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-3-chloro-N-[(1-methylethoxy)-
carbonyl]phenylalanine;

4-[[[3-[[[aminoiminomethyl)amino]carbonyl]phenyl]-
carbonyl]amino]-N-[(1-methylethoxy)-
carbonyl]phenylalanine; and

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]- α -ethylbenzenepropanoic acid.

5. A compound according to Claim 2 wherein B is selected from the group consisting of $-\text{CH}_2\text{O}-$,



6. A compound according to Claim 5 wherein the compound is selected from the group consisting of

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4-[[3-[(aminoiminomethyl)amino]phenyl]-methoxy]-N-(butylsulfonyl)phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]-sulfonyl]amino]benzenepropanoic acid; and

N-acetyl-4-[[[3-[(aminoiminomethyl)amino]phenyl]amino]carbonyl]amino]phenylalanine.

7. A compound according to Claim 2 wherein B is selected from the group consisting of $-\text{CONR}^{52}(\text{CH}_2)_{1-2}-$ and $-\text{CH}_2\text{CONR}^{52}-$.

8. A compound according to Claim 7 wherein the compound is selected from the group consisting of

ethyl 4-[[[3-(aminoiminomethyl)phenyl]-carbonyl]amino]methyl]benzenepropanoate;

ethyl 4-[[[3-[amino(hydroxyimino)methyl]-phenyl]carbonyl]amino]methyl]benzenepropanoate;

4-[[[3-(aminoiminomethyl)phenyl]carbonyl]-amino]methyl]benzenepropanoic acid;

3-[[[3-[(aminocarbonyl)amino]phenyl]-carbonyl]amino]methyl]benzenepropanoic acid;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-amino]methyl]benzenepropanoic acid;

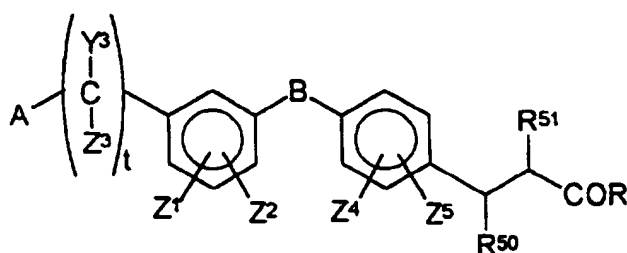
1,1-dimethylethyl-4-[2-[[3-[(aminoiminomethyl)-amino]phenyl]carbonyl]amino]ethyl]benzenepropanoate;

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4-[2-[[[3-[(aminoiminomethyl)amino]phenyl]-carbonyl]amino]ethyl]benzenepropanoic acid; and

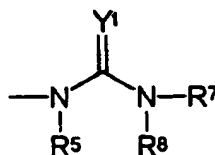
4-[[2-[4-[(aminoiminomethyl)amino]phenyl]-acetyl]amino]benzenepropanoic acid.

9. A pharmaceutical composition comprising a therapeutically effective amount of a compound of the formula



or a pharmaceutically acceptable salt thereof, wherein

A is



wherein Y^1 is selected from the group consisting of $N-R^2$, O, and S;

R^2 is selected from the group consisting of H; alkyl; aryl; hydroxy; alkoxy; cyano; nitro; amino; aminocarbonyl; alkenyl; alkynyl; alkyl optionally substituted with one or more substituent selected from lower alkyl, halogen, hydroxyl, haloalkyl, cyano, nitro, carboxyl, amino, alkoxy, aryl or aryl optionally substituted with one or more halogen,

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haloalkyl, lower alkyl, alkoxy, cyano, alkylsulfonyl, alkylthio, nitro, carboxyl, amino, hydroxyl, sulfonic acid, sulfonamide, aryl, fused aryl, monocyclic heterocycles, or fused monocyclic heterocycles; aryl optionally substituted with one or more substituent selected from halogen, haloalkyl, hydroxy, lower alkyl, alkoxy, methylenedioxy, ethylenedioxy, cyano, nitro, alkylthio, alkylsulfonyl, sulfonic acid, sulfonamide, carboxyl derivatives, amino, aryl, fused aryl, monocyclic heterocycles and fused monocyclic heterocycle; monocyclic heterocycles; and monocyclic heterocycles optionally substituted with one or more substituent selected from halogen, haloalkyl, lower alkyl, alkoxy, amino, nitro, hydroxy, carboxyl derivatives, cyano, alkylthio, alkylsulfonyl, sulfonic acid, sulfonamide, aryl or fused aryl; or

R^2 taken together with R^7 forms a 4-12 membered dinitrogen containing heterocycle optionally substituted with one or more substituent selected from the group consisting of lower alkyl, hydroxy and phenyl and wherein said ring optionally contains a heteroatom selected from the group consisting of O and S;

or R^2 taken together with R^7 forms a 5 membered heteroaromatic ring or optionally substituted with amino;

or R^2 taken together with R^7 forms a 5 membered heteroaromatic ring fused with a phenyl group;

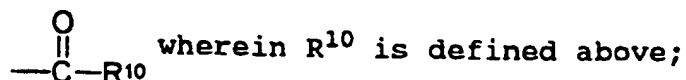
R^7 (when not taken together with R^2) and R^8 are independently selected from the group consisting of

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H; alkyl; alkenyl; alkynyl; aralkyl; cycloalkyl; bicycloalkyl; aryl; acyl; benzoyl; alkyl optionally substituted with one or more substituent selected from lower alkyl, halogen, hydroxy, haloalkyl, cyano, nitro, carboxyl derivatives, amino, alkoxy, thio, alkylthio, sulfonyl, aryl, aralkyl, aryl optionally substituted with one or more substituent selected from halogen, haloalkyl, lower alkyl, alkoxy, methylenedioxy, ethylenedioxy, alkylthio, haloalkylthio, thio, hydroxy, cyano, nitro, carboxyl derivatives, aryloxy, amido, acylamino, amino, alkylamino, dialkylamino, trifluoroalkoxy, trifluoromethyl, sulfonyl, alkylsulfonyl, haloalkylsulfonyl, sulfonic acid, sulfonamide, aryl, fused aryl, monocyclic heterocycles, fused monocyclic heterocycles; aryl optionally substituted with one or more substituent selected from halogen, haloalkyl, lower alkyl, alkoxy, methylenedioxy, ethylenedioxy, alkylthio, haloalkylthio, thio, hydroxy, cyano, nitro, carboxyl derivatives, aryloxy, amido, acylamino, amino, alkylamino, dialkylamino, trifluoroalkoxy, trifluoromethylsulfonyl, alkylsulfonyl, sulfonic acid, sulfonamide, aryl, fused aryl, monocyclic heterocycles, or fused monocyclic heterocycles; monocyclic heterocycles; monocyclic heterocycles optionally substituted with one or more substituent selected from halogen, haloalkyl, lower alkyl, alkoxy, aryloxy, amino, nitro, hydroxy, carboxyl derivatives, cyano, alkylthio, alkylsulfonyl, aryl, fused aryl; monocyclic and bicyclic heterocyclicalalkyls; $-SO_2R^{10}$ wherein R^{10} is selected from the group consisting of alkyl, aryl and monocyclic heterocycles, all optionally substituted with one or more substituent

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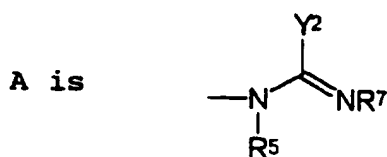
selected from the group consisting of halogen, haloalkyl, alkyl, alkoxy, cyano, nitro, amino, acylamino, trifluoroalkyl, amido, alkylaminosulfonyl, alkylsulfonyl, alkylsulfonylamino, alkylamino, dialkylamino, trifluoromethylthio, trifluoroalkoxy, trifluoromethylsulfonyl, aryl, aryloxy, thio, alkylthio, and monocyclic heterocycles; and



or NR^7 and R^8 taken together form a 4-12 membered mononitrogen containing monocyclic or bicyclic ring optionally substituted with one or more substituent selected from lower alkyl, carboxyl derivatives, aryl or hydroxy and wherein said ring optionally contains a heteroatom selected from the group consisting of O, N and S;

R^5 is selected from the group consisting of H, alkyl, alkenyl, alkynyl, benzyl, and phenethyl;

or



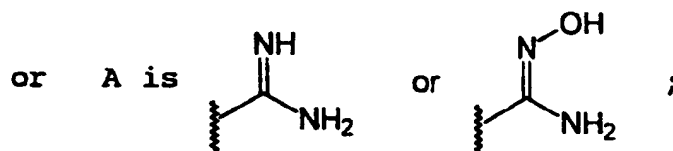
wherein Y^2 is selected from the group consisting of H, alkyl; cycloalkyl; bicycloalkyl; aryl; monocyclic heterocycles; alkyl optionally substituted with aryl which can also be optionally substituted with one or more substituent selected from halo, haloalkyl,

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alkyl, nitro, hydroxy, alkoxy, aryloxy, aryl, or fused aryl; aryl optionally substituted with one or more substituent selected from halo, haloalkyl, hydroxy, alkoxy, aryloxy, aryl, fused aryl, nitro, methylenedioxy, ethylenedioxy, or alkyl; alkynyl; alkenyl; $-S-R^9$ and $-O-R^9$ wherein R^9 is selected from the group consisting of H; alkyl; aralkyl; aryl; alkenyl; and alkynyl; or R^9 taken together with R^7 forms a 4-12 membered mononitrogen containing sulfur or oxygen containing heterocyclic ring; and

R^5 and R^7 are as defined above;

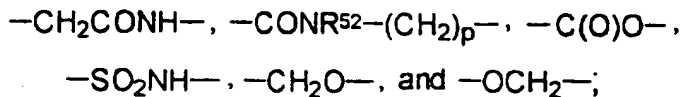
or Y^2 (when Y^2 is carbon) taken together with R^7 forms a 4-12 membered mononitrogen containing ring optionally substituted with alkyl, aryl or hydroxy;



Z^1 , Z^2 , Z^4 and Z^5 are independently selected from the group consisting of H; alkyl; hydroxy; alkoxy; aryloxy; aralkoxy; halogen; haloalkyl; haloalkoxy; nitro; amino; aminoalkyl; alkylamino; dialkylamino; cyano; alkylthio; alkylsulfonyl; carboxyl derivatives; carboxyalkyl; alkoxy-carbonylalkyl; acetamide; aryl; fused aryl; cycloalkyl; thio; monocyclic heterocycles; fused monocyclic heterocycles; and A, wherein A is defined above;

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B is selected from the group consisting of

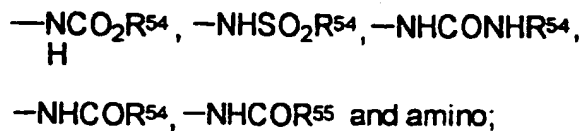


wherein p is an integer selected from the group consisting of 0, 1 and 2;

wherein R^{50} is selected from the group consisting of H, alkyl, aryl, carboxyl derivative and - $\text{CONHCH}_2\text{CO}_2\text{R}^{53}$ wherein R^{53} is H or lower alkyl;

R^{52} is selected from the group consisting of H or alkyl;

R^{51} is selected from the group consisting of H, alkyl, carboxyl derivatives,



wherein R^{54} is selected from the group consisting of H, alkyl, cycloalkyl, aryl, aralkyl aralkenyl and aryl substituted by one or more alkyl or halo;
wherein R^{55} is selected from the group consisting of N-substituted pyrrolidinyl, piperidinyl and morpholinyl;

t is an integer 0, 1 or 2;

R is X-R^3 wherein X is selected from the group consisting of O, S and NR^4 , wherein R^3 and R^4 are independently selected from the group consisting of

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hydrogen; alkyl; alkenyl; alkynyl; haloalkyl; aryl; arylalkyl; sugars; steroids and in the case of the free acid, all pharmaceutically acceptable salts thereof;

Y^3 and Z^3 are independently selected from the group consisting of H, alkyl, aryl, cycloalkyl and aralkyl or Y^3 and Z^3 taken together with C form a carbonyl;

and a pharmaceutically acceptable carrier.

10. A pharmaceutical composition according to Claim 9 wherein t is O.
11. A pharmaceutical composition according to Claim 10 wherein B is $-\text{CONR}^{52}-$.
12. A pharmaceutical composition according to Claim 11 selected from the group consisting of

4-[[[3-[(aminoiminomethyl)amino]-5-(trifluoromethyl)-phenyl]carbonyl]amino]-N-[(2-methylpropoxy)-carbonyl]-L-phenylalanine, methyl ester;

(\pm) 4-[[[3-[(aminoiminomethyl)amino]-5-(trifluoromethyl)phenyl]carbonyl]amino]-N-[(2-methylpropoxy)carbonyl]phenylalanine, methyl ester;

N-[(2-methylpropoxy)carbonyl]-4-[[[3-[(3,4,5,6-tetrahydro-2H-azepin-7-yl)amino]-5-(trifluoromethyl)phenyl]carbonyl]amino]-L-phenylalanine, methyl ester;

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N-[(2-methylpropoxy)carbonyl]-
4-[[[3-[(3,4,5,6-tetrahydro-2H-azepin-7-yl)-
amino]-5-(trifluoromethyl)phenyl]carbonyl]-
amino]phenylalanine;

N-(butylsulfonyl)-4-[[[3-
[(3,4,5,6-tetrahydro-2H-azepin-7-yl) amino]-5-
(trifluoromethyl)phenyl]carbonyl]amino]-
phenylalanine, methyl ester;

N-(butylsulfonyl)-
4-[[[3-[(3,4,5,6-tetrahydro-2H-azepin-7-yl)-
amino]-5-(trifluoromethyl)phenyl]carbonyl]-
amino]phenylalanine;

4-[[[3-(aminoiminomethyl)amino]-
5-(trifluoromethyl)phenyl]carbonyl]-
amino]-N-(butylsulfonyl)-phenylalanine,
methyl ester;

4-[[[3-(aminoiminomethyl)amino]-
5-(trifluoromethyl)phenyl]carbonyl]-
amino]-N-(butylsulfonyl)phenylalanine;

N-(butylsulfonyl)-4-[[[3-[(3,4,5,6-tetrahydro-
2H-azepin-7-yl) amino]-phenyl]carbonyl]amino] /
-phenylalanine, methyl ester;

N-(butylsulfonyl)-4-[[[3-[(3,4,5,6-
tetrahydro-2H-azepin-7-yl) amino]phenyl]-
carbonyl]amino]phenylalanine;

ethyl 4-[[[3-[amino(hydroxyimino)methyl]-
phenyl]carbonyl]amino]benzenepropanoate;

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ethyl 3-[[[3-(aminoiminomethyl)phenyl]-
carbonyl]amino]methyl]benzenepropanoate;

ethyl 3-[[[3-(aminoiminomethyl)phenyl]-
carbonyl]amino]benzenepropanoate;

4-[[[3-(aminoiminomethyl)phenyl]carbonyl]-
amino]benzenepropanoic acid;

ethyl 4-[[[3-(aminocarbonyl)amino]phenyl]-
carbonyl]amino]benzenepropanoate;

4-[[[3-(aminocarbonyl)amino]phenyl]-
carbonyl]amino]benzenepropanoic acid;

ethyl 4-[[[3-[[[(phenylmethyl)amino]carbonyl]-
amino]phenyl]carbonyl]amino]benzenepropanoate;

4-[[[3-[[[(phenylmethyl)amino]carbonyl]amino]-
phenyl]carbonyl]amino]benzenepropanoic acid;

1,1-dimethylethyl 4-[[[3-(aminoiminomethyl)-
amino]phenyl]carbonyl]amino]- β -
methylbenzenepropanoate;

4-[[[3-(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]- β -methylbenzenepropanoic acid;

1,1-dimethylethyl 4-[[[3-(aminoiminomethyl)-
amino]phenyl]carbonyl]amino]- β -
phenyl benzenepropanoate;

4-[[[3-(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]- β -phenylbenzenepropanoic acid;

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N-acetyl-4-[[[3-[(aminoiminomethyl)amino]-phenyl]carbonyl]amino]phenylalanine ethyl ester;

N-acetyl-4-[[[3-[(aminoiminomethyl)amino]-phenyl]carbonyl]amino]phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-amino]-N-(phenylcarbonyl)phenylalanine ethyl ester;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-amino]-N-(methylsulfonyl)phenylalanine ethyl ester;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-amino]-N-(ethoxycarbonyl)phenylalanine ethyl ester;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-amino]-N-[(1-methylethoxy)carbonyl]-phenylalanine ethyl ester;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-amino]-N-(phenylsulfonyl)phenylalanine ethyl ester;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-amino]-N-(butylsulfonyl)phenylalanine ethyl ester;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-amino]-N-[(2-methylpropoxy)carbonyl]-phenylalanine, ethyl ester;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-amino]-N-[(phenoxy)carbonyl]phenylalanine, ethyl ester;

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4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(1,1-dimethylethoxy)carbonyl]-
phenylalanine ethyl ester;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(1,1-dimethylethoxy)carbonyl]-
phenylalanine, methyl ester;

4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]amino]-N-[(1,1-dimethylethoxy)-
carbonyl]-L-phenylalanine, ethyl ester;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-(phenylcarbonyl)phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-(methylsulfonyl)phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-(ethoxycarbonyl)phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(1-methylethoxy)carbonyl]phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-(phenylsulfonyl)phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-(butylsulfonyl)phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(2-methylpropoxy)carbonyl]phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-(methoxycarbonyl)phenylalanine;

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4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]amino]-
N-[(1,1-dimethylethoxy)carbonyl]-phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(1,1-dimethylethoxy)carbonyl]-
L-phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(1,1-dimethylethoxy)carbonyl]-
phenylalanine;

ethyl 4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]methylamino]benzenepropanoate;

4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]methylamino]benzenepropanoic acid;

ethyl 4-[[[3-[(aminoiminomethyl)amino]-
phenyl]carbonyl]amino]benzenepropanoate;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]benzenepropanoic acid, hydrochloride;

4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]amino]phenylalanine;

[4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]amino]phenyl]butanedioic acid;

4-(1,1-dimethylethyl) 1-ethyl[4-[[[3-
[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]phenyl]butanedioate;

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1-ethyl hydrogen[4-[[[3-[(aminoiminomethyl)amino]-phenyl]carbonyl]amino]-phenyl]butanedioate;

1,1-dimethylethyl 4-[[[3-[(aminoiminomethyl)amino]-phenyl]carbonyl]-amino]- β -[[[(ethoxycarbonyl)-methyl]amino]-carbonyl]benzenepropanoate;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-amino]-beta-[(carboxymethyl)amino]-carbonyl]-benzenepropanoic acid;

1,1-dimethylethyl 4-[[[3-[(aminoiminomethyl)amino]-phenyl]carbonyl]amino]-3-hydroxybenzenepropanoate;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-amino]-3-hydroxybenzenepropanoic acid;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-amino]-N-[(1,1-dimethylethoxy)carbonyl]-2-methoxyphenylalanine, ethyl ester;

4-[[[3-[(4,5-dihydro-4-oxo-1H-imidazol-2-yl)amino]-phenyl]carbonyl]amino]-N-[(1-methylethoxy)-carbonyl]phenylalanine;

4-[[[3-[[[(cyanoimino)[(phenylmethyl)-amino]methyl]amino]phenyl]carbonyl]amino]-benzenepropanoic acid;

[4-[2-[[3-[(aminoiminomethyl)amino]phenyl]-amino]-2-oxoethyl]phenoxy]acetic acid;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-carbonyl]amino]benzene-1,2-dipropanoic acid;

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4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-2-carboxybenzenepropanoic acid;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-2-methoxycarbonyl)benzenepropanoic acid;

2-carboxy-4-[[[3-[(1,4,5,6-tetrahydropyrimidin-
2-yl)amino]phenyl]carbonyl]amino]-
benzenepropanoic acid;

(±) 4-[[[3-[3-amino-1H-1,2,4-triazol-5-yl)amino]-
phenyl]carbonyl]amino]-N-[(1-methylethoxy)
carbonyl]phenylalanine;

(±) 4-[[[3-[(5-amino-1,2,4-oxadiazol-3-yl)amino]-
phenyl]carbonyl]amino]-N-[(1-methylethoxy)
carbonyl]phenylalanine;

4-[[[3-[[amino[(aminocarbonyl)imino]methyl]-
amino]phenyl]carbonyl]amino]-N-[(1-methylethoxy)-
carbonyl]phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(3-methyl-1-oxobutyl)phenylalanine;

N-(3-methyl-1-oxobutyl)-4-[[[3-[(1,4,5,6-tetrahydro-
pyrimidin-2-yl)amino]-5-(trifluoromethyl)phenyl]-
carbonyl]amino]phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]-5-
(trifluoromethyl)phenyl]carbonyl]amino]-
N-(3-methyl-1-oxobutyl)phenylalanine;

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4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-(3,3-dimethyl-1-oxobutyl)phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]-5-(trifluoromethyl)-
phenyl]carbonyl]amino]-N-(3,3-dimethyl-
1-oxobutyl)phenylalanine;

N-(phenylsulfonyl)-4-[[[3-[(1,4,5,6-tetrahydro-
pyrimidin-2-yl)amino]phenyl]carbonyl]-
amino]phenylalanine;

4-[[[3-[(4,5-dihydro-1H-imidazol-2-yl)amino]phenyl]-
carbonyl]amino]-N-(phenylsulfonyl)phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]amino]-
2-methoxy-N-[(1-methylethoxy)carbonyl]phenylalanine;

2-methoxy-N-[(1-methylethoxy)carbonyl]-4-
[[[3-[(1,4,5,6-tetrahydropyrimidin-2-yl)amino]-
phenyl]carbonyl]amino]phenylalanine;

4-[[[3-(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-2-methoxybenzenepropanoic acid;

2-methoxy-4-[[[3-[(1,4,5,6-tetrahydropyrimidin-2-yl)-
amino]phenyl]carbonyl]amino]benzenepropanoic acid;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-2-(dimethylamino)benzenepropanoic acid;

[4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-2-methoxyphenoxy]acetic acid;

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[2-methoxy-4-[[[3-[(1,4,5,6-tetrahydropyrimidin-2-yl)-amino]phenyl]carbonyl]amino]phenoxy]acetic acid;

[[4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-amino]phenyl]methyl]-1,3-propanoic acid;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-amino]-N-[(2,4,6-trimethylphenyl)-sulfonyl]phenylalanine;

[4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]amino]-N-[(3,5-dichlorophenyl)sulfonyl]phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]amino]-N-[(2-phenylethenyl)sulfonyl]phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-amino]-N-[(1,1-dimethylethyl)amino]-carbonyl]phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-amino]-N-[(4-morpholinyl)carbonyl]phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-amino]-N-[(1-pipridinyl)carbonyl]phenylalanine;

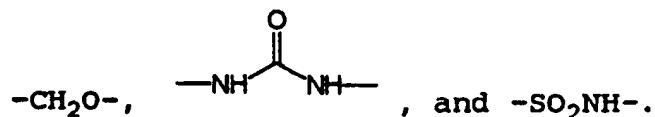
4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-amino]-3-chloro-N-[(1-methylethoxy)-carbonyl]phenylalanine;

4-[[[3-[[[3-[(aminoiminomethyl)amino]carbonyl]phenyl]-carbonyl]amino]-N-[(1-methylethoxy)-carbonyl]phenylalanine; and

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4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-amino]- α -ethylbenzenepropanoic acid.

13. A pharmaceutical composition according to Claim 10 wherein B is selected from the group consisting of



14. A pharmaceutical composition according to Claim 13 wherein the compound is selected from the group consisting of

4-[[3-[(aminoiminomethyl)amino]phenyl]-methoxy]-N-(butylsulfonyl)phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]-sulfonyl]amino]benzenepropanoic acid; and

N-acetyl-4-[[[3-[(aminoiminomethyl)amino]phenyl]amino]carbonyl]amino]phenylalanine.

15. A pharmaceutical composition according to Claim 10 wherein B is selected from the group consisting of $-\text{CONR}^{52}(\text{CH}_2)_{1-2}-$ and $-\text{CH}_2\text{CONR}^{52}-$.

16. A pharmaceutical composition according to Claim 15 wherein the compound is selected from the group consisting of

ethyl 4-[[[3-(aminoiminomethyl)phenyl]-carbonyl]amino]methyl]benzenepropanoate;

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ethyl 4-[[[3-[amino(hydroxyimino)methyl]-phenyl]carbonyl]amino]methyl]benzenepropanoate;

4-[[[3-(aminoiminomethyl)phenyl]carbonyl]-amino]methyl]benzenepropanoic acid;

3-[[[3-[(aminocarbonyl)amino]phenyl]-carbonyl]amino]methyl]benzenepropanoic acid;

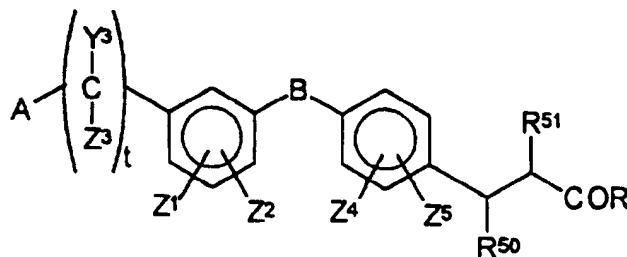
4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-amino]methyl]benzenepropanoic acid;

1,1-dimethylethyl-4-[2-[[[3-[(aminoiminomethyl)-amino]phenyl]carbonyl]amino]ethyl]benzenepropanoate;

4-[2-[[[3-[(aminoiminomethyl)amino]phenyl]-carbonyl]amino]ethyl]benzenepropanoic acid; and

4-[[2-[4-[(aminoiminomethyl)amino]phenyl]-acetyl]amino]benzenepropanoic acid.

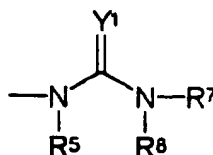
17. A method for treating conditions mediated by the $\alpha_v\beta_3$ integrin in a mammal in need of such treatment comprising administering a therapeutically effective $\alpha_v\beta_3$ inhibiting amount of a compound of the formula



or a pharmaceutically acceptable salt thereof, wherein

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A is



wherein Y^1 is selected from the group consisting of N- R^2 , O, and S;

R^2 is selected from the group consisting of H; alkyl; aryl; hydroxy; alkoxy; cyano; nitro; amino; aminocarbonyl; alkenyl; alkynyl; alkyl optionally substituted with one or more substituent selected from lower alkyl, halogen, hydroxyl, haloalkyl, cyano, nitro, carboxyl, amino, alkoxy, aryl or aryl optionally substituted with one or more halogen, haloalkyl, lower alkyl, alkoxy, cyano, alkylsulfonyl, alkylthio, nitro, carboxyl, amino, hydroxyl, sulfonic acid, sulfonamide, aryl, fused aryl, monocyclic heterocycles, or fused monocyclic heterocycles; aryl optionally substituted with one or more substituent selected from halogen, haloalkyl, hydroxy, lower alkyl, alkoxy, methylenedioxy, ethylenedioxy, cyano, nitro, alkylthio, alkylsulfonyl, sulfonic acid, sulfonamide, carboxyl derivatives, amino, aryl, fused aryl, monocyclic heterocycles and fused monocyclic heterocycle; monocyclic heterocycles; and monocyclic heterocycles optionally substituted with one or more substituent selected from halogen, haloalkyl, lower alkyl, alkoxy, amino, nitro, hydroxy, carboxyl derivatives, cyano, alkylthio, alkylsulfonyl, sulfonic acid, sulfonamide, aryl or fused aryl; or

R^2 taken together with R^7 forms a 4-12 membered dinitrogen containing heterocycle optionally substituted with one or more substituent selected from

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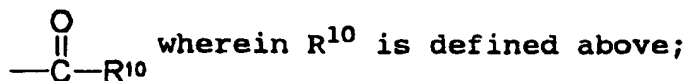
the group consisting of lower alkyl, hydroxy and phenyl and wherein said ring optionally contains a heteroatom selected from the group consisting of O and S;

- or R^2 taken together with R^7 forms a 5 membered heteroaromatic ring optionally substituted by amino;
- or R^2 taken together with R^7 forms a 5 membered heteroaromatic ring fused with a phenyl group;

R^7 (when not taken together with R^2) and R^8 are independently selected from the group consisting of H; alkyl; alkenyl; alkynyl; aralkyl; cycloalkyl; bicycloalkyl; aryl; acyl; benzoyl; alkyl optionally substituted with one or more substituent selected from lower alkyl, halogen, hydroxy, haloalkyl, cyano, nitro, carboxyl derivatives, amino, alkoxy, thio, alkylthio, sulfonyl, aryl, aralkyl, aryl optionally substituted with one or more substituent selected from halogen, haloalkyl, lower alkyl, alkoxy, methylenedioxy, ethylenedioxy, alkylthio, haloalkylthio, thio, hydroxy, cyano, nitro, carboxyl derivatives, aryloxy, amido, acylamino, amino, alkylamino, dialkylamino, trifluoroalkoxy, trifluoromethyl, sulfonyl, alkylsulfonyl, haloalkylsulfonyl, sulfonic acid, sulfonamide, aryl, fused aryl, monocyclic heterocycles, fused monocyclic heterocycles; aryl optionally substituted with one or more substituent selected from halogen, haloalkyl, lower alkyl, alkoxy, methylenedioxy, ethylenedioxy, alkylthio, haloalkylthio, thio, hydroxy, cyano, nitro, carboxyl derivatives, aryloxy, amido, acylamino, amino, alkylamino, dialkylamino, trifluoroalkoxy, trifluoromethylsulfonyl, alkylsulfonyl, sulfonic acid,

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sulfonamide, aryl, fused aryl, monocyclic heterocycles, or fused monocyclic heterocycles; monocyclic heterocycles; monocyclic heterocycles optionally substituted with one or more substituent selected from halogen, haloalkyl, lower alkyl, alkoxy, aryloxy, amino, nitro, hydroxy, carboxyl derivatives, cyano, alkylthio, alkylsulfonyl, aryl, fused aryl; monocyclic and bicyclic heterocyclicalkyls; $-\text{SO}_2\text{R}^{10}$ wherein R^{10} is selected from the group consisting of alkyl, aryl and monocyclic heterocycles, all optionally substituted with one or more substituent selected from the group consisting of halogen, haloalkyl, alkyl, alkoxy, cyano, nitro, amino, acylamino, trifluoroalkyl, amido, alkylaminosulfonyl, alkylsulfonyl, alkylsulfonylamino, alkylamino, dialkylamino, trifluoromethylthio, trifluoroalkoxy, trifluoromethylsulfonyl, aryl, aryloxy, thio, alkylthio, and monocyclic heterocycles; and



or NR^7 and R^8 taken together form a 4-12 membered mononitrogen containing monocyclic or bicyclic ring optionally substituted with one or more substituent selected from lower alkyl, carboxyl derivatives, aryl or hydroxy and wherein said ring optionally contains a heteroatom selected from the group consisting of O, N and S;

R^5 is selected from the group consisting of H, alkyl, alkenyl, alkynyl, benzyl, and phenethyl;

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N-(butylsulfonyl)-4-[[[3-[(3,4,5,6-tetrahydro-2H-azepin-7-yl)amino]phenyl]-carbonyl]amino]phenylalanine;

4-[[3-[(aminoiminomethyl)amino]phenyl]-methoxy]-N-(butylsulfonyl)phenylalanine;

ethyl 4-[[[3-(aminoiminomethyl)phenyl]-carbonyl]amino]methyl]benzenepropanoate;

ethyl 4-[[[3-[amino(hydroxyimino)methyl]-phenyl]carbonyl]amino]methyl]benzenepropanoate;

ethyl 4-[[[3-[amino(hydroxyimino)methyl]-phenyl]carbonyl]amino]benzenepropanoate;

ethyl 3-[[[3-(aminoiminomethyl)phenyl]-carbonyl]amino]methyl]benzenepropanoate;

ethyl 3-[[[3-(aminoiminomethyl)phenyl]-carbonyl]amino]benzenepropanoate;

4-[[[3-(aminoiminomethyl)phenyl]carbonyl]-amino]methyl]benzenepropanoic acid;

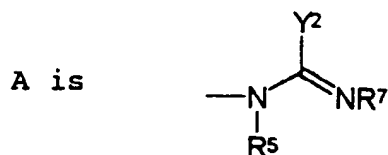
4-[[[3-(aminoiminomethyl)phenyl]carbonyl]-amino]benzenepropanoic acid;

ethyl 4-[[[3-[(aminocarbonyl)amino]phenyl]-carbonyl]amino]benzenepropanoate;

3-[[[3-[(aminocarbonyl)amino]phenyl]-carbonyl]amino]methyl]benzenepropanoic acid;

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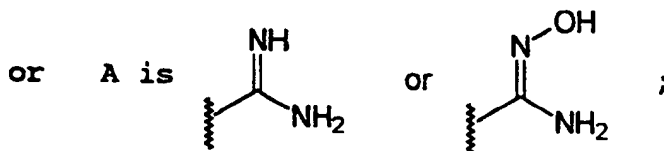
or



wherein Y^2 is selected from the group consisting of H, alkyl; cycloalkyl; bicycloalkyl; aryl; monocyclic heterocycles; alkyl optionally substituted with aryl which can also be optionally substituted with one or more substituent selected from halo, haloalkyl, alkyl, nitro, hydroxy, alkoxy, aryloxy, aryl, or fused aryl; aryl optionally substituted with one or more substituent selected from halo, haloalkyl, hydroxy, alkoxy, aryloxy, aryl, fused aryl, nitro, methylenedioxy, ethylenedioxy, or alkyl; alkynyl; alkenyl; $-\text{S}-\text{R}^9$ and $-\text{O}-\text{R}^9$ wherein R^9 is selected from the group consisting of H; alkyl; aralkyl; aryl; alkenyl; and alkynyl; or R^9 taken together with R^7 forms a 4-12 membered mononitrogen containing sulfur or oxygen containing heterocyclic ring; and

R^5 and R^7 are as defined above;

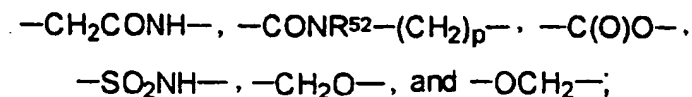
or Y^2 (when Y^2 is carbon) taken together with R^7 forms a 4-12 membered mononitrogen containing ring optionally substituted with alkyl, aryl or hydroxy;



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Z^1 , Z^2 , Z^4 and Z^5 are independently selected from the group consisting of H; alkyl; hydroxy; alkoxy; aryloxy; aralkoxy; halogen; haloalkyl; haloalkoxy; nitro; amino; aminoalkyl; alkylamino; dialkylamino; cyano; alkylthio; alkylsulfonyl; carboxyl derivatives; carboxyalkyl; alkoxy-carbonylalkyl; acetamide; aryl; fused aryl; cycloalkyl; thio; monocyclic heterocycles; fused monocyclic heterocycles; and A, wherein A is defined above;

B is selected from the group consisting of

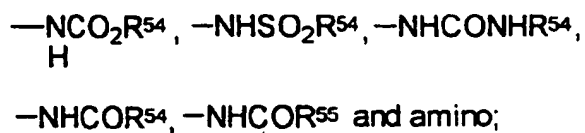


wherein p is an integer selected from the group consisting of 0, 1 and 2;

wherein R^{50} is selected from the group consisting of H, alkyl, aryl, carboxyl derivative and $-\text{CONHCH}_2\text{CO}_2R^{53}$ wherein R^{53} is H or lower alkyl;

R^{52} is selected from the group consisting of H or alkyl;

R^{51} is selected from the group consisting of H, alkyl, carboxyl derivatives,



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wherein R^{54} is selected from the group consisting of H, alkyl, cycloalkyl, aryl, aralkyl, aralkenyl and aryl substituted by one or more alkyl or halo; wherein R^{55} is selected from the group consisting of N-substituted pyrrolidinyl, piperidinyl and morpholinyl;

t is an integer 0, 1 or 2;

R is $X-R^3$ wherein X is selected from the group consisting of O, S and NR^4 , wherein R^3 and R^4 are independently selected from the group consisting of hydrogen; alkyl; alkenyl; alkynyl; haloalkyl; aryl; arylalkyl; sugars; steroids and in the case of the free acid, all pharmaceutically acceptable salts thereof; and

Y^3 and Z^3 are independently selected from the group consisting of H, alkyl, aryl, cycloalkyl and aralkyl or Y^3 and Z^3 taken together with C form a carbonyl.

18. The method according to Claim 17 wherein the compound is selected from the group consisting of

4-[[[3-[(aminoiminomethyl)amino]-5-(trifluoromethyl)-phenyl]carbonyl]amino]-N-[(2-methylpropoxy)-carbonyl]-phenylalanine, methyl ester;

4-[[[3-[(aminoiminomethyl)amino]-5-(trifluoromethyl)phenyl]carbonyl]amino]-N-[(2-methylpropoxy)carbonyl]phenylalanine, methyl ester;

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N-[(2-methylpropoxy)carbonyl]-4-[[[3-
[(3,4,5,6-tetrahydro-2H-azepin-7-yl)amino]-5-
(trifluoromethyl)phenyl]carbonyl]amino]-
phenylalanine, methyl ester;

N-[(2-methylpropoxy)carbonyl]-
4-[[[3-[(3,4,5,6-tetrahydro-2H-azepin-7-yl)-
amino]-5-(trifluoromethyl)phenyl]carbonyl]-
amino]phenylalanine;

N-(butylsulfonyl)-4-[[[3-
[(3,4,5,6-tetrahydro-2H-azepin-7-yl)amino]-5-
(trifluoromethyl)phenyl]carbonyl]amino]-
phenylalanine, methyl ester;

N-(butylsulfonyl)-4-[[[3-[(3,4,5,6-tetrahydro-
2H-azepin-7-yl)amino]-5-(trifluoromethyl)-
phenyl]carbonyl]amino]phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]-
5-(trifluoromethyl)phenyl]carbonyl]-
amino]-N-(butylsulfonyl)-phenylalanine,
methyl ester;

4-[[[3-[(aminoiminomethyl)amino]-
5-(trifluoromethyl)phenyl]carbonyl]-
amino]-N-(butylsulfonyl)phenylalanine;

N-(butylsulfonyl)-4-[[[3-[(3,4,5,6-tetrahydro-
2H-azepin-7-yl)amino]-phenyl]carbonyl]amino]-
phenylalanine, methyl ester;

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4-[[[3-[(aminocarbonyl)amino]phenyl]-
carbonyl]amino]benzenepropanoic acid;

ethyl 4-[[[3-[[[(phenylmethyl)amino]carbonyl]-
amino]phenyl]carbonyl]amino]benzenepropanoate;

4-[[[3-[[[(phenylmethyl)amino]carbonyl]amino]-
phenyl]carbonyl]amino]benzenepropanoic acid;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]methyl]benzenepropanoic acid;

1,1-dimethylethyl 4-[[[3-[(aminoiminomethyl)-
amino]phenyl]carbonyl]amino]- β -
methylbenzenepropanoate;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]- β -methylbenzenepropanoic acid;

1,1-dimethylethyl 4-[[[3-[(aminoiminomethyl)-
amino]phenyl]carbonyl]amino] β -
phenyl benzenepropanoate;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]- β -phenylbenzenepropanoic acid;

N-acetyl-4-[[[3-[(aminoiminomethyl)amino]-
phenyl]carbonyl]amino]phenylalanine ethyl ester;

N-acetyl-4-[[[3-[(aminoiminomethyl)amino]-
phenyl]carbonyl]amino]phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-(phenylcarbonyl)phenylalanine ethyl ester;

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4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-(methylsulfonyl)phenylalanine ethyl ester;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-(ethoxycarbonyl)phenylalanine ethyl ester;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(1-methylethoxy)carbonyl]-
phenylalanine ethyl ester;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-(phenylsulfonyl)phenylalanine ethyl ester;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-(butylsulfonyl)phenylalanine ethyl ester;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(2-methylpropoxy)carbonyl]-
phenylalanine, ethyl ester;

4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]amino]-N-[(phenoxy)carbonyl]-
phenylalanine, ethyl ester;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(1,1-dimethylethoxy)carbonyl]-
phenylalanine ethyl ester;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(1,1-dimethylethoxy)carbonyl]-
phenylalanine, methyl ester;

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- 4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]amino]-N-[(1,1-dimethylethoxy)-
carbonyl]-L-phenylalanine, ethyl ester;
- 4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-(phenylcarbonyl)phenylalanine;
- 4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-(methylsulfonyl)phenylalanine;
- 4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-(ethoxycarbonyl)phenylalanine;
- 4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(1-methylethoxy)carbonyl]phenylalanine;
- 4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-(phenylsulfonyl)phenylalanine;
- 4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-(butylsulfonyl)phenylalanine;
- 4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(2-methylpropoxy)carbonyl]phenylalanine;
- 4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-(methoxycarbonyl)phenylalanine;
- 4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]amino]-N-[(1,1-dimethylethoxy)-
carbonyl]-phenylalanine;

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4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(1,1-dimethylethoxy)carbonyl]-
L-phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(1,1-dimethylethoxy)carbonyl]-
phenylalanine;

1,1-dimethylethyl-4-[2-[[[3-[(aminoiminomethyl)-
amino]phenyl]carbonyl]amino]ethyl]benzenepropanoate;

4-[2-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]amino]ethyl]benzenepropanoic acid;

ethyl 4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]methylamino]benzenepropanoate;

4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]methylamino]benzenepropanoic acid;

4-[[[3-[(aminoiminomethyl)amino]phenyl]-
sulfonyl]amino]benzenepropanoic acid;

ethyl 4-[[[3-[(aminoiminomethyl)amino]-
phenyl]carbonyl]amino]benzenepropanoate;

4-[[[3-[(aminoiminomethyl)amino]phenyl]
carbonyl]amino]benzenepropanoic acid;

4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]amino]phenylalanine;

N-acetyl-4-[[[3-[(aminoiminomethyl)amino]-
phenyl]amino]carbonyl]amino]phenylalanine;

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[4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]amino]phenyl]butanedioic acid;

4-(1,1-dimethylethyl) 1-ethyl[4-[[[3-
[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]phenyl]butanedioate;

1-ethyl hydrogen[4-[[[3-[(aminoiminomethyl)amino]-
phenyl]carbonyl]amino]-phenyl]butanedioate;

1,1-dimethylethyl 4-[[[3-[(aminoiminomethyl)amino]-
phenyl]carbonyl]-amino]- β -[[[(ethoxycarbonyl)-
methyl]amino]-carbonyl]benzenepropanoate;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-beta-[[[carboxymethyl]amino]-
carbonyl]-benzenepropanoic acid;

1,1-dimethylethyl 4-[[[3-[(aminoiminomethyl)amino]-
phenyl]carbonyl]amino]-3-hydroxybenzenepropanoate;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-3-hydroxybenzenepropanoic acid;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(1,1-dimethylethoxy)carbonyl]-
2-methoxyphenylalanine, ethyl ester;

4-[[[3-[(4,5-dihydro-4-oxo-1H-imidazol-2-yl)amino]-
phenyl]carbonyl]amino]-N-[(1-methylethoxy)-
carbonyl]phenylalanine;

4-[[[2-[4-[(aminoiminomethyl)amino]phenyl]-
acetyl]amino]benzenepropanoic acid;

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4-[[[3-[(cyanoimino)(phenylmethyl)-amino]methyl]amino]phenyl]carbonyl]amino]-benzenepropanoic acid;

[4-[2-[[3-(aminoiminomethyl)amino]phenyl]-amino]-2-oxoethyl]phenoxy]acetic acid;

4-[[[3-(aminoiminomethyl)amino]phenyl]carbonyl]-carbonyl]amino]benzene-1,2-dipropanoic acid;

4-[[[3-(aminoiminomethyl)amino]phenyl]carbonyl]-amino]-2-carboxybenzenepropanoic acid;

4-[[[3-(aminoiminomethyl)amino]phenyl]carbonyl]-amino]-2-methoxycarbonyl]benzenepropanoic acid;

2-carboxy-4-[[[3-[(1,4,5,6-tetrahydropyrimidin-2-yl)amino]phenyl]carbonyl]amino]-benzenepropanoic acid;

(±) 4-[[[3-[3-amino-1H-1,2,4-triazol-5-yl]amino]phenyl]carbonyl]amino]-N-(1-methylethoxy)carbonyl]phenylalanine;

(±) 4-[[[3-[(5-amino-1,2,4-oxadiazol-3-yl)amino]phenyl]carbonyl]amino]-N-(1-methylethoxy)carbonyl]phenylalanine;

4-[[[3-[amino(aminocarbonyl)imino]methyl]-amino]phenyl]carbonyl]amino]-N-(1-methylethoxy)carbonyl]phenylalanine;

4-[[[3-(aminoiminomethyl)amino]phenyl]carbonyl]-amino]-N-(3-methyl-1-oxobutyl)phenylalanine;

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N-(3-methyl-1-oxobutyl)-4-[[[3-[(1,4,5,6-tetrahydro-
pyrimidin-2-yl)amino]-5-(trifluoromethyl)phenyl]-
carbonyl]amino]phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]-5-
(trifluoromethyl)phenyl]carbonyl]amino]-
N-(3-methyl-1-oxobutyl)phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-(3,3-dimethyl-1-oxobutyl)phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]-5-(trifluoromethyl)-
phenyl]carbonyl]amino]-N-(3,3-dimethyl-
1-oxobutyl)phenylalanine;

N-(phenylsulfonyl)-4-[[[3-[(1,4,5,6-tetrahydro-
pyrimidin-2-yl)amino]phenyl]carbonyl]-
amino]phenylalanine;

4-[[[3-[(4,5-dihydro-1H-imidazol-2-yl)amino]phenyl]-
carbonyl]amino]-N-(phenylsulfonyl)phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]amino]-
2-methoxy-N-[(1-methylethoxy)carbonyl]phenylalanine;

2-methoxy-N-[(1-methylethoxy)carbonyl]-4-
[[[3-[(1,4,5,6-tetrahydropyrimidin-2-yl)amino]-
phenyl]carbonyl]amino]phenylalanine;

4-[[[3-(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-2-methoxybenzenepropanoic acid;

2-methoxy-4-[[[3-[(1,4,5,6-tetrahydropyrimidin-2-yl)-
amino]phenyl]carbonyl]amino]benzenepropanoic acid;

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4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-2-(dimethylamino)benzenepropanoic acid;

[4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-2-methoxyphenoxy]acetic acid;

[2-methoxy-4-[[[3-[(1,4,5,6-tetrahydropyrimidin-2-yl)-
amino]phenyl]carbonyl]amino]phenoxy]acetic acid;

[[4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]phenyl]methyl]-1,3-propanoic acid;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(2,4,6-trimethylphenyl)-
sulfonyl]phenylalanine;

[4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]amino]-
N-[(3,5-dichlorophenyl)sulfonyl]phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]amino]-
N-[(2-phenylethenyl)sulfonyl]phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[[[(1,1-dimethylethyl)amino]-
carbonyl]phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(4-morpholinyl)carbonyl]phenylalanine;

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-N-[(1-pipecridinyl)carbonyl]phenylalanine;

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4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]-3-chloro-N-[(1-methylethoxy)-
carbonyl]phenylalanine;

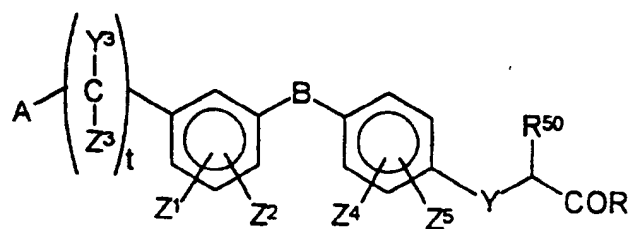
4-[[[3-[(aminoiminomethyl)amino]carbonyl]phenyl]-
carbonyl]amino]-N-[(1-methylethoxy)-
carbonyl]phenylalanine; and

4-[[[3-[(aminoiminomethyl)amino]phenyl]carbonyl]-
amino]- α -ethylbenzenepropanoic acid.

19. A method according to Claim 17 wherein the condition treated is tumor metastasis.
20. A method according to Claim 18 wherein the condition treated is tumor metastasis.
21. A method according to Claim 17 wherein the condition treated is solid tumor growth.
22. A method according to Claim 18 wherein the condition treated is solid tumor growth.
23. A method according to Claim 17 wherein the condition treated is angiogenesis.
24. A method according to Claim 18 wherein the condition treated is angiogenesis.
25. A method according to Claim 17 wherein the condition treated is osteoporosis.
26. A method according to Claim 18 wherein the condition treated is osteoporosis.

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27. A method according to Claim 17 wherein the condition treated is humoral hypercalcemia of malignancy.
28. A method according to Claim 18 wherein the condition treated is humoral hypercalcemia of malignancy.
29. A method according to Claim 17 wherein the condition treated is smooth muscle cell migration.
30. A method according to Claim 18 wherein the condition treated is smooth muscle cell migration.
31. A method according to Claim 17 wherein restenosis is inhibited.
32. A method according to Claim 18 wherein restenosis is inhibited.
33. A compound of the formula

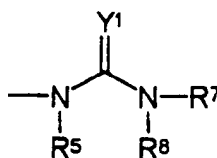


or a pharmaceutically acceptable salt thereof, wherein

Y is selected from the group consisting of -O-; -S-;
and -SO₂;

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A is



wherein Y¹ is selected from the group consisting of N-R², O, and S;

R² is selected from the group consisting of H; alkyl; aryl; hydroxy; alkoxy; cyano; nitro; amino; aminocarbonyl; alkenyl; alkynyl; alkyl optionally substituted with one or more substituent selected from lower alkyl, halogen, hydroxyl, haloalkyl, cyano, nitro, carboxyl, amino, alkoxy, aryl or aryl optionally substituted with one or more halogen, haloalkyl, lower alkyl, alkoxy, cyano, alkylsulfonyl, alkylthio, nitro, carboxyl, amino, hydroxyl, sulfonic acid, sulfonamide, aryl, fused aryl, monocyclic heterocycles, or fused monocyclic heterocycles; aryl optionally substituted with one or more substituent selected from halogen, haloalkyl, hydroxy, lower alkyl, alkoxy, methylenedioxy, ethylenedioxy, cyano, nitro, alkylthio, alkylsulfonyl, sulfonic acid, sulfonamide, carboxyl derivatives, amino, aryl, fused aryl, monocyclic heterocycles and fused monocyclic heterocycle; monocyclic heterocycles; and monocyclic heterocycles optionally substituted with one or more substituent selected from halogen, haloalkyl, lower alkyl, alkoxy, amino, nitro, hydroxy, carboxyl derivatives, cyano, alkylthio, alkylsulfonyl, sulfonic acid, sulfonamide, aryl or fused aryl; or

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R^2 taken together with R^7 forms a 4-12 membered dinitrogen containing heterocycle optionally substituted with one or more substituent selected from the group consisting of lower alkyl, hydroxy and phenyl and wherein said ring optionally contains a heteroatom selected from the group consisting of O and S;

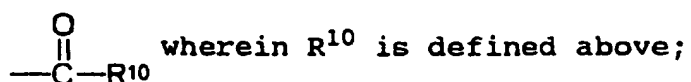
or R^2 taken together with R^7 forms a 5 membered heteroaromatic ring optionally substituted by amino;

or R^2 taken together with R^7 forms a 5 membered heteroaromatic ring fused with a phenyl group;

R^7 (when not taken together with R^2) and R^8 are independently selected from the group consisting of H; alkyl; alkenyl; alkynyl; aralkyl; cycloalkyl; bicycloalkyl; aryl; acyl; benzoyl; alkyl optionally substituted with one or more substituent selected from lower alkyl, halogen, hydroxy, haloalkyl, cyano, nitro, carboxyl derivatives, amino, alkoxy, thio, alkylthio, sulfonyl, aryl, aralkyl, aryl optionally substituted with one or more substituent selected from halogen, haloalkyl, lower alkyl, alkoxy, methylenedioxy, ethylenedioxy, alkylthio, haloalkylthio, thio, hydroxy, cyano, nitro, carboxyl derivatives, aryloxy, amido, acylamino, amino, alkylamino, dialkylamino, trifluoroalkoxy, trifluoromethyl, sulfonyl, alkylsulfonyl, haloalkylsulfonyl, sulfonic acid, sulfonamide, aryl, fused aryl, monocyclic heterocycles, fused monocyclic heterocycles; aryl optionally substituted with one or more substituent selected from halogen, haloalkyl, lower alkyl, alkoxy, methylenedioxy, ethylenedioxy, alkylthio, haloalkylthio, thio, hydroxy, cyano, nitro, carboxyl

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derivatives, aryloxy, amido, acylamino, amino, alkylamino, dialkylamino, trifluoroalkoxy, trifluoromethylsulfonyl, alkylsulfonyl, sulfonic acid, sulfonamide, aryl, fused aryl, monocyclic heterocycles, or fused monocyclic heterocycles; monocyclic heterocycles; monocyclic heterocycles optionally substituted with one or more substituent selected from halogen, haloalkyl, lower alkyl, alkoxy, aryloxy, amino, nitro, hydroxy, carboxyl derivatives, cyano, alkylthio, alkylsulfonyl, aryl, fused aryl; monocyclic and bicyclic heterocyclicalkyls; $-\text{SO}_2\text{R}^{10}$ wherein R^{10} is selected from the group consisting of alkyl, aryl and monocyclic heterocycles, all optionally substituted with one or more substituent selected from the group consisting of halogen, haloalkyl, alkyl, alkoxy, cyano, nitro, amino, acylamino, trifluoroalkyl, amido, alkylaminosulfonyl, alkylsulfonyl, alkylsulfonylamino, alkylamino, dialkylamino, trifluoromethylthio, trifluoroalkoxy, trifluoromethylsulfonyl, aryl, aryloxy, thio, alkylthio, and monocyclic heterocycles; and

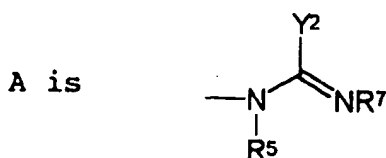


or NR^7 and R^8 taken together form a 4-12 membered mononitrogen containing monocyclic or bicyclic ring optionally substituted with one or more substituent selected from lower alkyl, carboxyl derivatives, aryl or hydroxy and wherein said ring optionally contains a heteroatom selected from the group consisting of O, N and S;

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R⁵ is selected from the group consisting of H, alkyl, alkenyl, alkynyl, benzyl, and phenethyl;

or

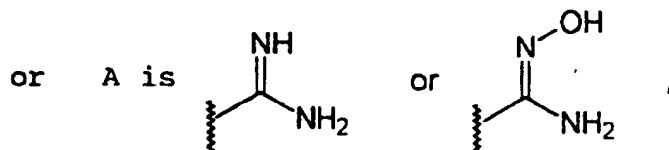


wherein Y² is selected from the group consisting of alkyl; cycloalkyl; bicycloalkyl; aryl; monocyclic heterocycles; alkyl optionally substituted with aryl which can also be optionally substituted with one or more substituent selected from halo, haloalkyl, alkyl, nitro, hydroxy, alkoxy, aryloxy, aryl, or fused aryl; aryl optionally substituted with one or more substituent selected from halo, haloalkyl, hydroxy, alkoxy, aryloxy, aryl, fused aryl, nitro, methylenedioxy, ethylenedioxy, or alkyl; alkynyl; alkenyl; -S-R⁹ and -O-R⁹ wherein R⁹ is selected from the group consisting of H; alkyl; aralkyl; aryl; alkenyl; and alkynyl; or R⁹ taken together with R⁷ forms a 4-12 membered mononitrogen containing sulfur or oxygen containing heterocyclic ring; and

R⁵ and R⁷ are as defined above;

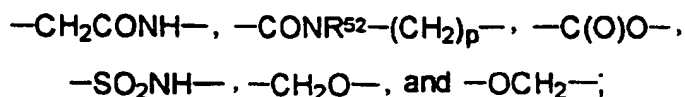
or Y² (when Y² is carbon) taken together with R⁷ forms a 4-12 membered mononitrogen containing ring optionally substituted with alkyl, aryl or hydroxy;

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Z^1 , Z^2 , Z^4 and Z^5 are independently selected from the group consisting of H; alkyl; hydroxy; alkoxy; aryloxy; aralkoxy; halogen; haloalkyl; haloalkoxy; nitro; amino; aminoalkyl; alkylamino; dialkylamino; cyano; alkylthio; alkylsulfonyl; carboxyl derivatives; carboxyalkyl; alkoxy-carbonylalkyl; acetamide; aryl; fused aryl; cycloalkyl; thio; monocyclic heterocycles; fused monocyclic heterocycles; and A, wherein A is defined above;

B is selected from the group consisting of



wherein p is an integer selected from the group consisting of 0, 1 and 2;

wherein R^{50} is selected from the group consisting of H, alkyl, aryl and carboxyl derivative;

R^{52} is selected from the group consisting of H or alkyl;

t is an integer 0, 1 or 2;

R is $\text{X}-\text{R}^3$ wherein X is selected from the group consisting of O, S and NR^4 , wherein R^3 and R^4 are independently selected from the group consisting of

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hydrogen; alkyl; alkenyl; alkynyl; haloalkyl; aryl; arylalkyl; sugars; steroids and in the case of the free acid, all pharmaceutically acceptable salts thereof; and

Y^3 and Z^3 are independently selected from the group consisting of H, alkyl, aryl, cycloalkyl and aralkyl or Y^3 and Z^3 taken together with C form a carbonyl.

34. A compound according to Claim 33 wherein the compound is selected from the group consisting of

ethyl [[4-[[[3-[(aminoiminomethyl)amino]phenyl]-carbonyl]amino]phenyl]sulfonyl]acetate;

[4-[[[3-[(aminoiminomethyl)amino]phenyl]-carbonyl]amino]phenoxy]acetic acid;

[[4-[[[3-[(aminoiminomethyl)amino]phenyl]-carbonyl]amino]phenyl]thio]acetic acid;

[[4-[[[3-[(aminoiminomethyl)amino]phenyl]-carbonyl]amino]phenyl]sulfonyl]acetic acid;

[4-[2-[[3-[(aminoiminomethyl)amino]phenyl]-amino]-2-oxoethyl]phenoxy]acetic acid;

2-[4-[[[3-[(aminoiminomethyl)amino]phenyl]-carbonyl]amino]phenoxy]propanoic acid;

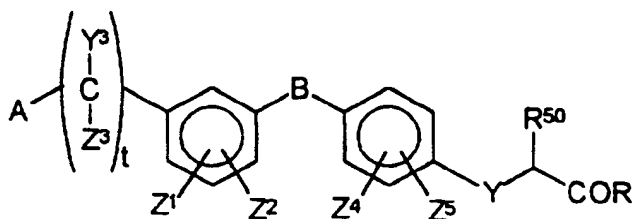
α -[4-[[[3-[(aminoiminomethyl)amino]phenyl]-carbonyl]amino]phenoxy]benzeneacetic acid;

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ethyl [4-[[[3-[(aminoiminomethyl)amino]-phenyl]carbonyl]amino]phenoxy]acetate; and

[[4-[[[3-[(aminoiminomethyl)amino]phenyl]-carbonyl]amino]phenyl]thio]acetic acid.

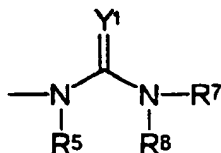
35. A pharmaceutical composition comprising a therapeutically effective amount of a compound of the formula



or a pharmaceutically acceptable salt thereof, wherein

Y is selected from the group consisting of -O-; -S-; and -SO₂;

A is



wherein Y¹ is selected from the group consisting of N-R², O, and S;

R² is selected from the group consisting of H; alkyl; aryl; hydroxy; alkoxy; cyano; nitro; amino; aminocarbonyl; alkenyl; alkynyl; alkyl optionally substituted with one or more substituent selected from lower alkyl, halogen, hydroxyl, haloalkyl, cyano, nitro, carboxyl, amino, alkoxy, aryl or aryl optionally

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substituted with one or more halogen, haloalkyl, lower alkyl, alkoxy, cyano, alkylsulfonyl, alkylthio, nitro, carboxyl, amino, hydroxyl, sulfonic acid, sulfonamide, aryl, fused aryl, monocyclic heterocycles, or fused monocyclic heterocycles; aryl optionally substituted with one or more substituent selected from halogen, haloalkyl, hydroxy, lower alkyl, alkoxy, methylenedioxy, ethylenedioxy, cyano, nitro, alkylthio, alkylsulfonyl, sulfonic acid, sulfonamide, carboxyl derivatives, amino, aryl, fused aryl, monocyclic heterocycles and fused monocyclic heterocycle; monocyclic heterocycles; and monocyclic heterocycles optionally substituted with one or more substituent selected from halogen, haloalkyl, lower alkyl, alkoxy, amino, nitro, hydroxy, carboxyl derivatives, cyano, alkylthio, alkylsulfonyl, sulfonic acid, sulfonamide, aryl or fused aryl; or

R^2 taken together with R^7 forms a 4-12 membered dinitrogen containing heterocycle optionally substituted with one or more substituent selected from the group consisting of lower alkyl, hydroxy and phenyl;

or R^2 taken together with R^7 forms a 5 membered heteroaromatic ring optionally substituted with amino;

or R^2 taken together with R^7 forms a 5 membered heteroaromatic ring fused with a phenyl group;

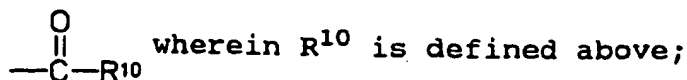
R^7 (when not taken together with R^2) and R^8 are independently selected from the group consisting of H; alkyl; alkenyl; alkynyl; aralkyl; cycloalkyl; bicycloalkyl; aryl; acyl; benzoyl; alkyl optionally

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substituted with one or more substituent selected from lower alkyl, halogen, hydroxy, haloalkyl, cyano, nitro, carboxyl derivatives, amino, alkoxy, thio, alkylthio, sulfonyl, aryl, aralkyl, aryl optionally substituted with one or more substituent selected from halogen, haloalkyl, lower alkyl, alkoxy, methylenedioxy, ethylenedioxy, alkylthio, haloalkylthio, thio, hydroxy, cyano, nitro, carboxyl derivatives, aryloxy, amido, acylamino, amino, alkylamino, dialkylamino, trifluoroalkoxy, trifluoromethyl, sulfonyl, alkylsulfonyl, haloalkylsulfonyl, sulfonic acid, sulfonamide, aryl, fused aryl, monocyclic heterocycles, fused monocyclic heterocycles; aryl optionally substituted with one or more substituent selected from halogen, haloalkyl, lower alkyl, alkoxy, methylenedioxy, ethylenedioxy, alkylthio, haloalkylthio, thio, hydroxy, cyano, nitro, carboxyl derivatives, aryloxy, amido, acylamino, amino, alkylamino, dialkylamino, trifluoroalkoxy, trifluoromethylsulfonyl, alkylsulfonyl, sulfonic acid, sulfonamide, aryl, fused aryl, monocyclic heterocycles, or fused monocyclic heterocycles; monocyclic heterocycles; monocyclic heterocycles optionally substituted with one or more substituent selected from halogen, haloalkyl, lower alkyl, alkoxy, aryloxy, amino, nitro, hydroxy, carboxyl derivatives, cyano, alkylthio, alkylsulfonyl, aryl, fused aryl; monocyclic and bicyclic heterocyclicalkyls; $-SO_2R^{10}$ wherein R^{10} is selected from the group consisting of alkyl, aryl and monocyclic heterocycles, all optionally substituted with one or more substituent selected from the group consisting of halogen, haloalkyl, alkyl, alkoxy, cyano, nitro, amino, acylamino, trifluoroalkyl, amido, alkylaminosulfonyl, alkylsulfonyl, alkylsulfonylamino,

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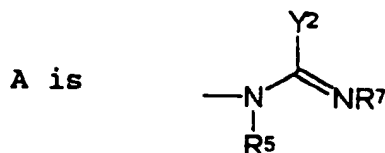
alkylamino, dialkylamino, trifluoromethylthio, trifluoroalkoxy, trifluoromethylsulfonyl, aryl, aryloxy, thio, alkylthio, and monocyclic heterocycles; and



or NR^7 and R^8 taken together form a 4-12 membered mononitrogen containing monocyclic or bicyclic ring optionally substituted with one or more substituent selected from lower alkyl, carboxyl derivatives, aryl or hydroxy and wherein said ring optionally contains a heteroatom selected from the group consisting of O, N and S;

R^5 is selected from the group consisting of H, alkyl, alkenyl, alkynyl, benzyl, and phenethyl;

or



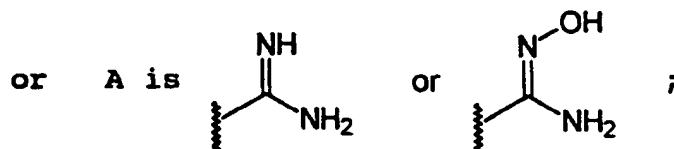
wherein Y^2 is selected from the group consisting of alkyl; cycloalkyl; bicycloalkyl; aryl; monocyclic heterocycles; alkyl optionally substituted with aryl which can also be optionally substituted with one or more substituent selected from halo, haloalkyl, alkyl, nitro, hydroxy, alkoxy, aryloxy, aryl, or fused aryl; aryl optionally substituted with one or more substituent selected from halo, haloalkyl, hydroxy,

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alkoxy, aryloxy, aryl, fused aryl, nitro, methylenedioxy, ethylenedioxy, or alkyl; alkynyl; alkenyl; -S-R⁹ and -O-R⁹ wherein R⁹ is selected from the group consisting of H; alkyl; aralkyl; aryl; alkenyl; and alkynyl; or R⁹ taken together with R⁷ forms a 4-12 membered mononitrogen containing sulfur or oxygen containing heterocyclic ring; and

R⁵ and R⁷ are as defined above;

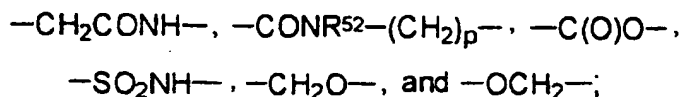
or Y² (when Y² is carbon) taken together with R⁷ forms a 4-12 membered mononitrogen containing ring optionally substituted with alkyl, aryl or hydroxy;



Z¹, Z², Z⁴ and Z⁵ are independently selected from the group consisting of H; alkyl; hydroxy; alkoxy; aryloxy; aralkoxy; halogen; haloalkyl; haloalkoxy; nitro; amino; aminoalkyl; alkylamino; dialkylamino; cyano; alkylthio; alkylsulfonyl; carboxyl derivatives; carboxyalkyl; alkoxycarbonylalkyl; acetamide; aryl; fused aryl; cycloalkyl; thio; monocyclic heterocycles; fused monocyclic heterocycles; and A, wherein A is defined above;

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B is selected from the group consisting of



wherein p is an integer selected from the group consisting of 0, 1 and 2;

wherein R^{50} is selected from the group consisting of H, alkyl, aryl and carboxyl derivative;

R^{52} is selected from the group consisting of H or alkyl;

t is an integer 0, 1 or 2;

R is X-R^3 wherein X is selected from the group consisting of O, S and NR^4 , wherein R^3 and R^4 are independently selected from the group consisting of hydrogen; alkyl; alkenyl; alkynyl; haloalkyl; aryl; arylalkyl; sugars; steroids and in the case of the free acid, all pharmaceutically acceptable salts thereof;

Y^3 and Z^3 are independently selected from the group consisting of H, alkyl, aryl, cycloalkyl and aralkyl or Y^3 and Z^3 taken together with C form a carbonyl;

and a pharmaceutically acceptable carrier.

36. A pharmaceutical composition according to Claim 35 wherein the compound is selected from the group consisting of

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ethyl [[4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]amino]phenyl]sulfonyl]acetate;

[4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]amino]phenoxy]acetic acid;

[[4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]amino]phenyl]thio]acetic acid;

[[4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]amino]phenyl]sulfonyl]acetic acid;

[4-[2-[3-[(aminoiminomethyl)amino]phenyl]-
amino]-2-oxoethyl]phenoxy]acetic acid;

2-[4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]amino]phenoxy]propanoic acid;

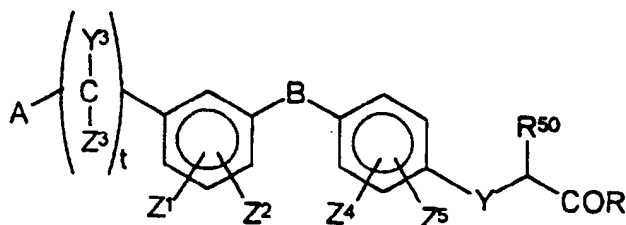
α -[4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]amino]phenoxy]benzeneacetic acid;

ethyl [4-[[[3-[(aminoiminomethyl)amino]-
phenyl]carbonyl]amino]phenoxy]acetate,
trifluoroacetate salt; and

[[4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]amino]phenyl]thio]acetic acid.

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37. A method for treating conditions mediated by the $\alpha_v\beta_3$ integrin in a mammal in need of such treatment comprising administering a therapeutically effective $\alpha_v\beta_3$ inhibiting amount of a compound of the formula



38. A method according to Claim 37 wherein the compound is selected from the group consisting of

ethyl [[4-[[[3-[(aminoiminomethyl)amino]phenyl]-carbonyl]amino]phenyl]sulfonyl]acetate;

[4-[[[3-[(aminoiminomethyl)amino]phenyl]-carbonyl]amino]phenoxy]acetic acid;

[[4-[[[3-[(aminoiminomethyl)amino]phenyl]-carbonyl]amino]phenyl]thio]acetic acid;

[[4-[[[3-[(aminoiminomethyl)amino]phenyl]-carbonyl]amino]phenyl]sulfonyl]acetic acid;

[4-[2-[[3-[(aminoiminomethyl)amino]phenyl]-amino]-2-oxoethyl]phenoxy]acetic acid;

2-[4-[[[3-[(aminoiminomethyl)amino]phenyl]-carbonyl]amino]phenoxy]propanoic acid;

α -[4-[[[3-[(aminoiminomethyl)amino]phenyl]-carbonyl]amino]phenoxy]benzeneacetic acid;

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ethyl [4-[[[3-[(aminoiminomethyl)amino]-
phenyl]carbonyl]amino]phenoxy]acetate; and

[[4-[[[3-[(aminoiminomethyl)amino]phenyl]-
carbonyl]amino]phenyl]thio]acetic acid.

39. A method according to Claim 37 wherein the condition treated is tumor metastasis.
40. A method according to Claim 38 wherein the condition treated is tumor metastasis.
41. A method according to Claim 37 wherein the condition treated is solid tumor growth.
42. A method according to Claim 38 wherein the condition treated is solid tumor growth.
43. A method according to Claim 37 wherein the condition treated is angiogenesis.
44. A method according to Claim 38 wherein the condition treated is angiogenesis.
45. A method according to Claim 37 wherein the condition treated is osteoporosis.
46. A method according to Claim 38 wherein the condition treated is osteoporosis.
47. A method according to Claim 37 wherein the condition treated is humoral hypercalcemia of malignancy.

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48. A method according to Claim 38 wherein the condition treated is humoral hypercalcemia of malignancy.
49. A method according to Claim 37 wherein the condition treated is smooth muscle cell migration.
50. A method according to Claim 38 wherein the condition treated is smooth muscle cell migration.
51. A method according to Claim 37 wherein restenosis is inhibited.
52. A method according to Claim 38 wherein restenosis is inhibited.

INTERNATIONAL SEARCH REPORT

International Application No
PC., US 97/04460

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C07C279/18 A61K31/155 C07D223/04 C07C311/06 C07C259/18
 C07C257/18 C07C275/42 C07C311/19 C07C311/47 C07D233/88
 C07C279/28 C07C317/44 C07C323/52 C07D239/14 C07D249/14

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C07C C07D A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
2	A EP 0 372 486 B (F. HOFFMANN-LA ROCHE) 1 June 1994 see page 6, line 7 - line 47; claims; examples ---	1-52
1	A WO 93 08823 A (TANABE SEIYAKU CO. LTD.) 13 May 1993 see claims; examples ---	1-52
1	A WO 92 08464 A (TANABE SEIYAKU CO. LTD.) 29 May 1992 see claims; examples ---	1-52
2	A US 5 352 667 A (LIDER ET. AL.) 4 October 1994 see claims; examples -----	1-52

☐ Further documents are listed in the continuation of box C.☒ Patent family members are listed in annex.

* Special categories of cited documents :

- "A" document defining the general state of the art which is not considered to be of particular relevance
 "E" earlier document but published on or after the international filing date
 "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
 "O" document referring to an oral disclosure, use, exhibition or other means
 "P" document published prior to the international filing date but later than the priority date claimed

- "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
 "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
 "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
 "&" document member of the same patent family

2

Date of the actual completion of the international search

31 July 1997

Date of mailing of the international search report

1 1. 08. 97

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Helps, I

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.: 17-32 & 37-52
because they relate to subject matter not required to be searched by this Authority, namely:

See extra sheet (A.)

2. ☒ Claims Nos.: 1-3, 5, 9-11, 13, 15, ...
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:

... 17, 19, 21, 23, 25, 27, 29, 31, 33, 35, 37, 39, 41, 43, 45, 47, 49, 51

See extra sheet (B.)

3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

☐ The additional search fees were accompanied by the applicant's protest.

☐ No protest accompanied the payment of additional search fees.